

From: McGarry, Sean
Sent: Wednesday, June 05, 2002 1:11 PM
To: STIC-Biotech/ChemLib
Subject: Sequence Search 09/599,220

Please,

For 09/599,220, a length limited search of SEQ ID NO: 1 and 2 (nucleotides < 50). Please do not search ESTs.

Thank You
Sean McGarry
AU 1635
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Office CM1 11D07
305-7028

Point of Contact:

Barb O'Bryen

Technical Information Specialist
STIC CM1 6A05 308-4291

Searcher: BOB
Phone: _____
Location: _____
Date Picked Up: _____
Date Completed: 6-7-02
Searcher Prep/Review: _____
Clerical: _____
Online time: _____

TYPE OF SEARCH:

NA Sequences: _____
AA Sequences: _____
Structures: _____
Bibliographic: _____
Litigation: _____
Full text: _____
Patent Family: _____
Other: _____

VENDOR/COST (where applic.)

STN: _____
DIALOG: _____
Questel/Orbit: _____
DRLink: _____
Lexis/Nexis: _____
Sequence Sys.: _____
WWW/Internet: _____
Other (specify): _____



Page
1

Copyright (c) 1993 - 2000 GenCore version OM nucleic - nucleic search, using sw model
Run on: June 6, 2002, 16:03:19 ; ;

GenCore version 4.5
copyright (c) 1993 - 2000 Compugen Ltd.
elc search, using sw model
June 6, 2002, 16:03:19 ; Search time 179

(without assignments)
337.739 Million cell updates/sec

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Searched:          1797656 seqs, 10463268293 residues
Scoring table:    IDENTITY.NUC
Post-processing:  Minimum Match 0%
                  Maximum Match 100%
                  Listing first 45 summaries
Database :        GenEmbl: *

```

Database : GenEmbl:*

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2: gb_htg:*
3: gb_in:*
4: gb_om:*
5: gb_out:*
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6:	gb.Bal:..	C	37	14.4
7:	gb.Ph:*			
8:	gb.pl:*		38	14.4
9:	gb.pr:*		39	14.2
10:	gb.ro:*	C	40	14.2
11:	gb.sts:*	C	41	14.2
12:	gb.sy:*	C	42	14.2
13:	gb.un:*	C	43	14.2
14:	gb.vi:*	C	44	14.2
15:	em.ba:*	C	45	14.2

ALIGNMENTS

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LOCUS	Sequence 279 from patent US 6177557.
DEFINITION	38 bp DNA
ACCESSION	AR125937
VERSION	AR125937.1 GI:14111999
KEYWORDS	
SOURCE	Unknown.
ORGANISM	Unclassified.
REFERENCE	1 (bases 1 to 38)
AUTHORS	Janik,N., Gold,L. and Tasset,D.
TITLE	High affinity ligands of basic fibroblast growth factor

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
------------	-------	-------------	--------	-------	-------------

BASE COUNT	6 a	6 c	17 g	10 t	RESULT	9	RESULT	9	RESULT	9	RESULT	9
ORIGIN					190181	190181	190181	190181	190181	190181	190181	190181
Query Match	100.0%	Score 29;	DB 6;	Length 39;	DEFINITION	Sequence 88 from patent US 5723594.	DEFINITION	Sequence 29;	DB 6;	Length 39;	DEFINITION	Sequence 88 from patent US 5723594.
Best Local Similarity	100.0%	Pred. No. 0.39;	Mismatches 0;	Indels 0;	ACCESSION	190181	ACCESSION	190181	ACCESSION	190181	ACCESSION	190181
Matches 29;	Conservative 0;	Mismatches 0;	Indels 0;	Gaps 0;	VERSION	190181.1	VERSION	190181.1	VERSION	190181.1	VERSION	190181.1
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Db	2	AGTCCGGTGGTAGGGCAGGTGGGTGACT 30			SOURCE	Unknown.	SOURCE	Unknown.	SOURCE	Unknown.	SOURCE	Unknown.
RESULT	7				REFERENCE	1 (bases 1 to 39)						
LOCUS	165730	165730	39 bp	DNA	ORGANISM	Janjic,N. and Gold,L.	ORGANISM	High affinity PDGF nucleic acid ligands	ORGANISM	High affinity PDGF nucleic acid ligands	ORGANISM	High affinity PDGF nucleic acid ligands
DEFINITION	Sequence 90 from patent US 5668264.			DEFINITION	Unclassified.	DEFINITION	Unclassified.	DEFINITION	Unclassified.	DEFINITION	Unclassified.	DEFINITION
ACCESSION	165730	165730.1	GI:2482300		VERSION	1.0	VERSION	1.0	VERSION	1.0	VERSION	1.0
VERSION					KEYWORDS		KEYWORDS		KEYWORDS		KEYWORDS	
KEYWORDS	.				SOURCE	Unknown.	SOURCE	Unknown.	SOURCE	Unknown.	SOURCE	Unknown.
REFERENCE					REFERENCE	1 (bases 1 to 39)						
AUTHORS	Janjic,N. and Gold,L.				ORGANISM	Unclassified.	ORGANISM	Unclassified.	ORGANISM	Unclassified.	ORGANISM	Unclassified.
TITLE	High affinity PDGF nucleic acid ligands				FEATURES		FEATURES		FEATURES		FEATURES	
JOURNAL	Patent: US 5668264-A 90 16-SEP-1997;				BASE COUNT	6 a						
FEATURES	Location/Qualifiers				ORGIN	6 c						
source	1. .39				RESULT	10	RESULT	10	RESULT	10	RESULT	10
BASE COUNT	6 a	6 c	17 g	10 t	LOCUS	AR125877	LOCUS	AR125877	LOCUS	AR125877	LOCUS	AR125877
ORIGIN					DEFINITION	Sequence 219 from patent US 6177557.	DEFINITION	Sequence 219 from patent US 6177557.	DEFINITION	Sequence 219 from patent US 6177557.	DEFINITION	Sequence 219 from patent US 6177557.
Query Match	100.0%	Score 29;	DB 6;	Length 39;	ACCESSION	AR125877	ACCESSION	AR125877	ACCESSION	AR125877	ACCESSION	AR125877
Best Local Similarity	100.0%	Pred. No. 0.39;	Mismatches 0;	Indels 0;	VERSION	AR125877.1	VERSION	AR125877.1	VERSION	AR125877.1	VERSION	AR125877.1
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Db	2	AGTCCGGTGGTAGGGCAGGTGGGTGACT 30			REFERENCE	1 (bases 1 to 30)						
RESULT	8				AUTHORS	Janjic,N. and Gold,L. and Tasset,D.						
LOCUS	167962	167962	39 bp	DNA	DEFINITION	High affinity ligands of basic fibroblast growth factor and	DEFINITION	High affinity ligands of basic fibroblast growth factor and	DEFINITION	High affinity ligands of basic fibroblast growth factor and	DEFINITION	High affinity ligands of basic fibroblast growth factor and
DEFINITION	Sequence 90 from patent US 5674685.			ACCESSION	167962	ACCESSION	167962	ACCESSION	167962	ACCESSION	167962	ACCESSION
ACCESSION	167962	167962.1	GI:2830084		VERSION	1.0	VERSION	1.0	VERSION	1.0	VERSION	1.0
VERSION					KEYWORDS		KEYWORDS		KEYWORDS		KEYWORDS	
KEYWORDS	.				SOURCE	Unclassified.	SOURCE	Unclassified.	SOURCE	Unclassified.	SOURCE	Unclassified.
REFERENCE					REFERENCE	1 (bases 1 to 30)						
AUTHORS	Janjic,N. and Gold,L.				TITLE	High affinity ligands of basic fibroblast growth factor and	TITLE	High affinity ligands of basic fibroblast growth factor and	TITLE	High affinity ligands of basic fibroblast growth factor and	TITLE	High affinity ligands of basic fibroblast growth factor and
TITLE	High affinity PDGF nucleic acid ligands				JOURNAL	Patent: US 6177557-A 219 23-JAN-2001;						
JOURNAL	Patent: US 5674685-A 90 07-OCT-1997;				FEATURES	Location/Qualifiers	FEATURES	Location/Qualifiers	FEATURES	Location/Qualifiers	FEATURES	Location/Qualifiers
FEATURES	Location/Qualifiers				source	1. .30						
source	1. .39	/organism="unknown"			BASE COUNT	5 a						
BASE COUNT	6 a	6 c	17 g	10 t	ORIGIN	14 g						
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Best Local Similarity	100.0%	Pred. No. 0.39;	Mismatches 0;	Indels 0;	DEFINITION	Sequence 31 from patent US 5543293.	DEFINITION	Sequence 31 from patent US 5543293.	DEFINITION	Sequence 31 from patent US 5543293.	DEFINITION	Sequence 31 from patent US 5543293.
Matches 29;	Conservative 0;	Mismatches 0;	Indels 0;	Gaps 0;	ACCESSION	124244	ACCESSION	124244	ACCESSION	124244	ACCESSION	124244
Qy	1	agtccgtggtagggcagggtgggtgact 29			VERSION	1.0	VERSION	1.0	VERSION	1.0	VERSION	1.0
Db	2	AGTCCGGTGGAGGCAGGTGGGTGACT 30			KEYWORDS	.	KEYWORDS	.	KEYWORDS	.	KEYWORDS	.
REFERENCE					SOURCE	Unknown.	SOURCE	Unknown.	SOURCE	Unknown.	SOURCE	Unknown.
ORGANISM	Unclassified.				REFERENCE	1 (bases 1 to 30)						
REFERENCE	1 (bases 1 to 30)				AUTHORS	Gold,L. and Tasset,D.						

Query	Match	Similarity	Score	DB	Length
Best	Local	95.2%	19.4	6	30
Matches	20;	Conservative	0;	Mismatches: 1;	Indels

Search completed: June 6, 2002, 16:03:21
Job time: 1822 sec

Gencore version 4.5
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OM nucleic - nucleic search, using SW model

Run on: June 6, 2002, 16:08:47 ; Search time 234.25 Seconds

(without alignments)
212.553 Million cell updates/sec

Title: US-09-599-220-2

Perfect score: 29

Sequence: 1 agtcgggttagggcagggtgggtact 29

Scoring table: IDENTITY_NUC

Gapop 10.0 , Gapext 1.0

Searched: 1736436 seqs, 858457221 residues

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

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Maximum DB seq length: 50

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2: /SIDS1/geodata/geneseq/geneseq/geneseq -emb1/NA1981 DAT:*

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15: /SIDS1/geodata/geneseq/geneseq -emb1/NA1994 DAT:*

16: /SIDS1/geodata/geneseq/geneseq -emb1/NA1995 DAT:*

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23: /SIDS1/geodata/geneseq/geneseq -emb1/NA2002 DAT:*

24: /SIDS1/geodata/geneseq/geneseq -emb1/NA2002 DAT:*

ALIGNMENTS

RESULT 1

ID AAC91745 standard; DNA; 29 BP.

XX AAC91745;

AC AAC91745;

DT 27-MAR-2001 (first entry)

XX DE Thrombin-binding aptamer. ODN 2.

XX KW Thrombin-binding aptamer; exosite 2; heparin binding site;

KW blood clot; anticoagulant; in vivo imaging; diagnostic tool;

KW protein quantitation; in vivo half-life; ss.

XX OS Synthetic.

XX PN WO200078364-A2.

XX PD 28-DEC-2000.

XX PF 22-JUN-2000; 2000WO-CA00751.

XX PR 22-JUN-1999; 99US-0139896.

XX PA (UYAL-) UNIV ALBERTA SIMON FRASER.

XX PI Dougan AH, Weitz JI;

XX DR WPI; 2001-091498/10.

XX PT Novel composition for inhibiting and preventing blood coagulation and

PT for imaging blood clots in vivo, comprises a nucleic acid that binds to

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query	Length	DB	ID	Description
1	29	100.0	29	22	ACG91745	Thrombin-binding a
2	29	100.0	37	24	AAS16530	Thrombin specific;
3	29	100.0	37	24	AAS16540	truncated S10EX de
4	29	100.0	38	16	AAU98404	Spectroscopically
5	29	100.0	38	17	AAU86654	Thrombin ligand fo
6	29	100.0	38	17	AAU38802	Thrombin-binding n
7	29	100.0	38	18	AAU85812	Thrombin binding l
8	29	100.0	38	18	AAU80049	Thrombin high affi
9	100.0	38	22	XX	AAU70817	

PR blood clot and complexed at its 5' or 3' end or both with a protein -
 XX
 PS Example 1; Page 5; 49pp; English.
 XX
 CC The invention relates to a composition comprising an aptamer which
 CC binds to a blood clot or to a protein component of a mammalian blood
 CC coagulation cascade, and a protein (other than the target protein)
 CC complexed with either or both the 5' and 3' ends of the aptamer. The
 CC proteins used in the composition can be covalently or non-covalently
 CC bound to the aptamer termini, and serve to extend the in vivo half-life
 CC of the aptamer. The aptamer/protein complexes of the invention are used
 CC to inhibit and prevent the coagulation of blood in a patient requiring
 CC anticoagulation treatment. Complexes may also be radiolabelled and used
 CC to image blood clots in vivo. The complexes can additionally be used to
 detect and quantitate the amount of a target protein in a sample to
 provide a diagnosis of a disease state that is correlated with the amount
 CC of protein in the sample. The present sequence represents an aptamer
 CC which binds to the heparin binding site (exosite 2) of human thrombin
 CC which was used in an exemplification of the invention.
 Sequence 29 BP; 4 A; 4 C; 14 G; 7 T; 0 other;

Query Match 100 0%; Score 29; DB 22; Length 29;
 Best Local Similarity 100 0%; Pred. No. 0.014; ID 16530; Mismatches 0; Indels 0; Gaps 0;
 Matches 29; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 Oy 1 agtcctgttagggagggtgggtgact 29
 DB 1 agtcctgttagggagggtgggtgact 29

RESULT 2
 ID AAS16530 standard; DNA; 37 BP.

XX
 AC AAS16530;
 XX
 DT 14-FEB-2002 (first entry)
 XX
 DE Thrombin specific, DT-aptamer.
 XX
 KW DT-aptamer; L-selectin; alpha-thrombin; plasma; blood;
 KW bronchial aspirate; sandwich assay; ss.
 XX
 OS Synthetic.
 XX

Key modified_base Location/Qualifiers
 FT /
 FT /
 FT misc_binding 30..37 13-12, 16-17 and 22-21 of this sequence*
 FT /*tag= 9
 FT /bound_moiety= "nucleotides 8-1"
 FT /note= "Forms a double stranded region with
 FT modified_base 37 nucleotides 8-1 of this sequence"
 FT /*tag= h
 FT /mod_base= c
 FT /note= " Optionally fluorescein labelled, if position 37
 FT is labelled, position 1 is not labelled"
 FT XX
 PN WO200179562-A1.
 PD 25-OCT-2001.
 PR 18-APR-2001; 2001WO-US12614.
 PR 18-APR-2000; 2000US-198016P.
 PA (GILE-) GILEAD SCI INC.
 XX
 PI Lin Y, Heil J, Jayasena S;
 DR XX
 PT WPI; 2002-017628/02.
 XX
 PT Novel aptamer based two-site binding sandwich assay for detecting
 target compounds such as thrombin and L-selectin in a biological fluid,
 PT employs nucleic acid ligands as capture and/or reporter molecules
 XX
 PS Example 1; Fig 1A; 47pp; English.

CC The invention describes a novel method of detecting the presence of a
 CC target compound in a substance which may contain the target compound. The
 CC method involves exposing the substance to a capture molecule (CM) capable
 CC of binding to the target molecule (TM) and immobilised on a solid
 CC support. A reporter molecule (RM) capable of binding to the target
 CC molecule is added to the CM:TM complex to detect the CM:TM:RM complex,
 CC where CM and/or RM are a nucleic acid ligand to TM. The method is useful
 CC for detecting a target molecule such as a protein, preferably thrombin or
 CC L-selectin in a biological fluid including plasma, blood and serum. The
 CC assays detect human alpha-thrombin in buffer as well as in biological
 CC fluids. Detection of the target compound is useful for clinical diagnosis
 CC of physiological conditions in both human and veterinary diagnostics. The
 CC nucleic acid ligand-based sandwich assays, designed on two different
 CC types of beads that can be readily analysed in flow cytometry, allow
 CC multiplexed analysis of a mixture of target protein in a single tube.
 CC This sequence is the alpha-thrombin specific DT-aptamer, the capture
 CC molecule used to detect alpha-thrombin in a sample using the method
 CC described in the invention.

XX
 SQ Sequence 37 BP; 6 A; 6 C; 16 G; 9 T; 0 other;

Query Match 100 0%; Score 29; DB 24; Length 37;
 Best Local Similarity 100 0%; Pred. No. 0.014; ID 16540; Mismatches 0; Indels 0; Gaps 0;
 Matches 29; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Oy 1 agtcctgttagggagggtgggtgact 29

DB 5 agtcctgttagggagggtgggtgact 33

RESULT 3
 ID AAS16540 standard; DNA; 37 BP.

XX
 AC AAS16540;
 XX
 DT 14-FEB-2002 (first entry)
 XX
 DE Thrombin specific, DT-DIMR-F aptamer.

FT misc_binding 30..37 13-12, 16-17 and 22-21 of this sequence*
 FT /*tag= 9
 FT /bound_moiety= "nucleotides 8-1"
 FT /note= "Forms a double stranded region with
 FT modified_base 37 nucleotides 8-1 of this sequence"
 FT /*tag= h
 FT /mod_base= c
 FT /note= " Optionally fluorescein labelled, if position 37
 FT is labelled, position 1 is not labelled"
 FT XX
 PN WO200179562-A1.
 PD 25-OCT-2001.
 PR 18-APR-2001; 2001WO-US12614.
 PR 18-APR-2000; 2000US-198016P.
 PA (GILE-) GILEAD SCI INC.
 XX
 PI Lin Y, Heil J, Jayasena S;
 DR XX
 PT WPI; 2002-017628/02.
 XX
 PT Novel aptamer based two-site binding sandwich assay for detecting
 target compounds such as thrombin and L-selectin in a biological fluid,
 PT employs nucleic acid ligands as capture and/or reporter molecules
 XX
 PS Example 1; Fig 1A; 47pp; English.

CC The invention describes a novel method of detecting the presence of a
 CC target compound in a substance which may contain the target compound. The
 CC method involves exposing the substance to a capture molecule (CM) capable
 CC of binding to the target molecule (TM) and immobilised on a solid
 CC support. A reporter molecule (RM) capable of binding to the target
 CC molecule is added to the CM:TM complex to detect the CM:TM:RM complex,
 CC where CM and/or RM are a nucleic acid ligand to TM. The method is useful
 CC for detecting a target molecule such as a protein, preferably thrombin or
 CC L-selectin in a biological fluid including plasma, blood and serum. The
 CC assays detect human alpha-thrombin in buffer as well as in biological
 CC fluids. Detection of the target compound is useful for clinical diagnosis
 CC of physiological conditions in both human and veterinary diagnostics. The
 CC nucleic acid ligand-based sandwich assays, designed on two different
 CC types of beads that can be readily analysed in flow cytometry, allow
 CC multiplexed analysis of a mixture of target protein in a single tube.
 CC This sequence is the alpha-thrombin specific DT-aptamer, the capture
 CC molecule used to detect alpha-thrombin in a sample using the method
 CC described in the invention.

XX
 SQ Sequence 37 BP; 6 A; 6 C; 16 G; 9 T; 0 other;

Query Match 100 0%; Score 29; DB 24; Length 37;
 Best Local Similarity 100 0%; Pred. No. 0.014; ID 16540; Mismatches 0; Indels 0; Gaps 0;
 Matches 29; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Oy 1 agtcctgttagggagggtgggtgact 29

DB 5 agtcctgttagggagggtgggtgact 33

RESULT 3
 ID AAS16540 standard; DNA; 37 BP.

XX
 AC AAS16540;
 XX
 DT 14-FEB-2002 (first entry)
 XX
 DE Thrombin specific, DT-DIMR-F aptamer.

DT-DIMR-F aptamer; L-selectin; alpha-thrombin; plasma; blood; KW
 KW bronchial aspirate; sandwich assay; ss.
 XX
 OS Synthetic.
 XX

KEY modified_base Location/Qualifiers
 FH 37
 FT /*tag^a a
 FT /mod_base^c
 FT /note= "two copies of this sequence are joined at the 3'
 FT end, by glycerol backbones, to a branching phosphoramidite is also
 FT phosphoramidite, the Phosphoramidite is also
 FT labelled by a phosphodester bond to a thymine
 FT labelled with fluorescein"
 XX
 PN WO2001179562-A1.
 XX
 PD 25-OCT-2001.
 XX
 PR 18-APR-2001; 2001WO-US12614.
 XX
 PR 18-APR-2000; 2000US-198016P.
 XX
 PA (GILE-.) GILEAD SCI INC.
 XX
 PI Lin Y, Heil J, Jayasena S;
 XX
 DR WPI; 2002-017628/02.
 XX
 PT Novel aptamer based two-site binding sandwich assay for detecting
 PT target compounds such as thrombin and L-selectin in a biological fluid,
 PT employs nucleic acid ligands as capture and/or reporter molecules.
 XX
 Disclosure: Page 32; 47pp; English.
 XX
 CC The invention describes a novel method of detecting the presence of a
 CC target compound in a substance which may contain the target compound. The
 CC method involves exposing the substance to a capture molecule (CM) capable
 CC of binding to the target molecule (TM) and immobilised on a solid
 CC support. A reporter molecule (RM) capable of binding to the target
 molecule is added to the CM:TM complex to detect the CM:TM:RM complex,
 CC where CM and/or RM are a nucleic acid ligand to TM. The method is useful
 CC for detecting a target molecule such as a protein, preferably thrombin or
 CC L-selectin in a biological fluid including plasma, blood and serum. The
 CC assays detect human alpha-thrombin in buffer as well as in biological
 CC fluids. Detection of the target compound is useful for clinical diagnosis
 CC of physiological conditions in both human and veterinary diagnostics. The
 CC nucleic acid ligand-based sandwich assays, designed on two different
 CC types of beads that can be readily analysed in flow cytometry, allow
 CC multiplexed analysis of a mixture of target protein in a single tube.
 CC This sequence is the alpha-thrombin specific DT-DIMR-F aptamer, a
 CC derivative of Dr-aptamer AS16530 consisting of two DR-aptamer joined to
 CC a fluorescein labelled branching phosphoramidite, this forms the capture
 CC molecule used to detect alpha-thrombin in a sample using the method
 CC described in the invention.
 XX
 Sequence 37 BP; 6 A; 6 C; 16 G; 9 T; 0 other;
 XX
 Query Match 100.0%; Score 29; DB 24; Length 37;
 Best Local Similarity 100.0%; Pred. No. 0.014; Indels 0; Gaps 0;
 Matches 29; Conservative 0; Mismatches 0;
 QY 1 agtccgtggatggcgggtgggtgact 29
 QY 2 agtccgtggatggcgggtgggtgact 30
 DB 5 agtccgtggatggcgggtgggtgact 33
 XX
 RESULT 4
 AAO98404
 ID AAO98404 standard; RNA; 38 BP.
 XX
 AC AAO98404;
 KW

XX
 DT 08-AUG-1996 (first entry)
 XX
 DE Truncated SELEX derived DNA thrombin ligand 60-18(38).
 XX
 KW Family 1; family 2; ligand; thrombin;
 KW systematic evolution of ligands by exponential enrichment; SELEX;
 KW heparin; selection; region of homology; inhibitor; ss.
 OS Synthetic.
 XX
 PN WO9521853-A1.
 XX
 PD 17-AUG-1995.
 XX
 PR 06-FEB-1995; 95WO-US01458.
 XX
 PR 28-MAR-1994; 94US-0219812.
 PR 10-FEB-1994; 94US-019505.
 PR 11-JUN-1990; 90US-0536428.
 PR 10-JUN-1991; 91US-0714131.
 PR 22-APR-1993; 93US-006191.
 XX
 PA (NEXS-) NEXSTAR PHARM INC.
 XX
 PI Gold L, Janjic N, Tasset D;
 XX
 DR WPI; 1995-293073/38.
 XX
 PT Identification of ligands to basic fibroblast growth factor and
 PT thrombin - which can be modified for increased in vivo stability
 XX
 PS Claim 39; Page 98; 236pp; English.
 XX
 CC The sequences given in AAO9837-405 represent DNA ligands directed to
 CC thrombin which were isolated using systematic evolution of ligands by
 CC exponential enrichment (SELEX). Two populations of single stranded
 CC (ss) DNA molecules with either 30N or 60N variable regions with 5', and
 CC 3', fixed regions were synthesised. Thrombin and DNA were incubated in
 CC a buffer at 37 deg.C for 5 mins. The thrombin-bound DNA is removed by
 CC filtration. A double stranded product was created and amplified by PCR,
 CC and a ssDNA template pool was isolated from this by alkaline
 CC denaturation. This ssDNA template pool was used for the following round
 CC of SELEX. Individual clones were isolated and the dissociation
 CC constants (Kd) were determined. Kd's ranged from 0.4-9.4 nM for the 30N
 CC ss's and from 0.9-2.5 nM for the 60N DNA's. A truncated ligand given in
 CC AAO9404 was derived from the high affinity clone 60-18 and has a Kd of
 CC 1.9 nM and inhibits clotting.
 XX
 SQ Sequence 38 BP; 6 A; 6 C; 17 G; 9 T; 0 other;
 XX
 Query Match 100.0%; Score 29; DB 16; Length 38;
 Best Local Similarity 100.0%; Pred. No. 0.014; Indels 0; Gaps 0;
 Matches 29; Conservative 0; Mismatches 0;
 QY 1 agtccgtggatggcgggtgggtgact 29
 QY 2 agtccgtggatggcgggtgggtgact 30
 DB
 RESULT 5
 AAT86654
 ID AAT86654 standard; DNA; 38 BP.
 XX
 AC AAT86654;
 XX
 DT 08-MAY-1998 (first entry)
 XX
 DE Spectroscopically detectable nucleic acid ligand compound #2.
 XX
 KW Spectroscopically detectable; detection; phosphorothioate;
 KW fluorescein; thiazole orange; ss.

linker arm (= Compound 2)"

linker arm (= Compound 2) "

```
misc_feature 1 /*tag= a /note= "fluorescein labelled"
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PD	22-JUL-1997.	PR	0550410423 R.
XX		XX	
PF	18-JUL-1994;	94US-0276271.	
XX		XX	
PF	18-JUN-1997.	PD	24-JUN-1997.
XX		XX	
PR	18-MAY-1995;	PR	11-JUN-1990;
PR	95US-0443357.	PR	90US-0536420.
PR	11-JUN-1990;	PR	18-MAY-1995;
PR	90US-0536428.	PR	95US-0443950.
PR	10-JUN-1991;	PR	11-JUN-1990;
PR	91US-0714131.	PR	90US-0536429.
PR	17-AUG-1992;	PR	10-JUN-1991;
PR	92US-0931473.	PR	91US-0714130.
PR	07-OCT-1993;	PR	17-AUG-1992;
PR	93US-0134028.	PR	92US-0931471.
PR	28-APR-1994;	PR	07-OCT-1993;
PR	94US-0234997.	PR	93US-0134022.
PR	18-JUL-1994;	PR	28-APR-1994;
XX	94US-0276271.	PR	94US-0244999.
PA	(GOLD/) GOLD L.	PR	18-JUL-1994;
PA	(MALIN/) MALINOWSKI D P.	PR	94US-0276270.
PA	(PTIN/) PITNER J B.	XX	95US-0376322.
PA	(YONK/) VONK G P.	PA	(GOLD/) GOLD L.
PA	(MALIN/) MALINOWSKI D P.	PA	

XX	Synthetic.
OS	
XX	
FH	
KEY	
	Location/Qualifiers

XX
 AC AAF70817;
 XX
 DT 20-APR-2001 (first entry)
 XX
 DE Thrombin high affinity ligand #64.
 XX
 KW Ligand; basic fibroblast growth factor; bFGF; gene therapy; vascular;
 XX
 KW atherosclerosis; angioplasty; stability; ss.
 XX
 OS Unidentified.
 XX
 PR US6177557-B1.
 XX
 PD 23-JAN-2001.
 XX
 PR 05-AUG-1996; 96US-0687421.
 XX
 PR 11-JUN-1990; 90US-0536428.
 XX
 PR 10-JUN-1991; 91US-0714131.
 XX
 PR 06-NOV-1992; 92US-0773333.
 XX
 PR 10-FEB-1994; 94US-0195005.
 XX
 PR 28-MAR-1994; 94US-0219012.
 XX
 PA (NEXS-) NEXSTAR PHARM INC.
 XX
 PI Janjic N, Gold L, Tasset D;
 XX
 DR WPI; 2001-158583/16.
 XX
 PT Novel nucleic acid ligands to basic fibroblast growth factor that are
 PT used as inhibitors of basic fibroblast growth factors and 2'-amino
 PT modified RNA ligands, exhibit increased in vivo stability.
 XX
 PS Example 19; column 61-62; 153pp; English.
 XX
 CC The present invention relates to a purified and isolated non-naturally
 CC occurring DNA ligands to basic fibroblast growth factor (bFGF).
 CC The ligands are useful as part of gene therapy treatments and
 CC for diagnosing pathogenesis of vascular diseases including
 CC initiation and progression of atherosclerosis, acute coronary
 CC syndromes, vein graft disease and restenosis following coronary
 CC angioplasty. The ligands have improved stability in vivo.
 XX
 SQ Sequence 38 BP; 6 A; 6 C; 17 G; 9 T; 0 other;
 XX
 SQ Query Match 100.0%; Score 29; DB 22; Length 38;
 XX
 Matches 29; Local Similarity 100.0%; Pred. No. 0.014; Indels 0; Gaps 0;
 XX
 QY 1 agtcccgtaggtggcagggtgggtact 29
 XX
 DB 2 agtcccgtaggtggcagggtgggtact 30
 XX
 RESULT 10
 ID AAX87088
 ID AAX87088 standard; DNA; 39 BP.
 XX
 AC AAX87088;
 XX
 DT 20-SEP-1999 (first entry)
 XX
 DE DNA ligand T39 to human thrombin.
 XX
 DE Platelet derived growth factor; PDGF; human; ligand; SELEX;
 KW systematic evolution of ligands by exponential enrichment;
 KW single stranded DNA; ssDNA; angiogenesis; restenosis; tumour;
 KW cancer; fibrosis; therapy; thrombin; ss.
 XX
 OS Synthetic.
 XX
 PR Synthetic.
 XX
 PA (NEXS-) NEXSTAR PHARM INC.
 XX
 PI Gold L, Janjic N;
 XX
 DR WPI; 1999-405022/34.
 XX
 PT Complex comprises a platelet derived growth factor nucleic acid
 PT ligand
 XX
 PS Example 3; Page 52; 156pp; English.
 XX
 CC This sequence represents DNA ligand T39 to human thrombin. It was
 CC used as a control in experiments designed to examine binding of
 CC minimal DNA ligands (see AAX8703-85) to human Platelet derived
 CC growth factor (PDGF). The invention discloses a method for
 CC preparing a complex of a PDGF nucleic acid ligand and a
 CC non-immunogenic high mol wt. compound (e.g., PEG) or lipophilic
 CC compound (e.g., a glycerol lipid). Such complexes are used as
 CC inhibitors of PDGF mediated angiogenesis, to inhibit the growth of
 CC tumours, to inhibit fibrosis (especially kidney, lung, bone marrow
 CC or radiation treatment associated fibrosis) or to inhibit
 CC restenosis, especially in-stent restenosis or restenosis in a
 CC coronary artery or non-coronary vessel. They can also be used to
 CC target a therapeutic or diagnostic agent to a biological target
 CC expressing PDGF.
 XX
 SQ Sequence 39 BP; 6 A; 6 C; 17 G; 10 T; 0 other;
 XX
 SQ Query Match 100.0%; Score 29; DB 20; Length 39;
 XX
 Matches 29; Local Similarity 100.0%; Pred. No. 0.014; Indels 0; Gaps 0;
 XX
 QY 1 agtcccgtaggtggcagggtgggtact 29
 XX
 DB 2 agtcccgtaggtggcagggtgggtact 30
 XX
 RESULT 11
 ID AAS16539
 ID AAS16539 standard; DNA; 42 BP.
 XX
 AC AAS16539;
 XX
 DT 14-FEB-2002 (first entry)
 XX
 DE Thrombin specific, Dn-5'-LNK aptamer.
 XX
 KW Dn-5'-LNK-aptamer; L-selectin; alpha-thrombin; plasma; blood;
 KW bronchial aspirate; sandwich assay; ss.
 XX
 OS Synthetic.
 XX
 PR Key
 PR modified_base
 PR 1
 PR /*tag= a
 PR /mod_base= t
 PR /label= Fluorescein
 XX

PI	LIN Y, HELL J, JAYASENA S;	XX
DR		XX
XX	Novel aptamer based two-site binding sandwich assay for detecting target compounds such as thrombin and L-selectin in a biological fluid.	XX
PT		XX
PT	Target compounds such as thrombin and L-selectin in a biological fluid.	XX
PS	Disclosure; Page 32; 47pp; English.	PS
XX	The invention describes a novel method of detecting the presence of a target compound in a substance which may contain the target compound. The method involves exposing the substance to a capture molecule (CM) capable of binding to the target molecule (TM) and immobilised on a solid support. A reporter molecule (RM) capable of binding to the target molecule is added to the CM:TM complex to detect the CM:TM:RM complex, where CM and/or RM are a nucleic acid ligand to TM. The method is useful for detecting a target molecule such as a protein, preferably thrombin or L-selectin in a biological fluid including plasma, blood and serum. The assays detect human alpha-thrombin in buffer as well as in biological fluids. Detection of the target compound is useful for clinical diagnosis of physiological conditions in both human and veterinary diagnostics. The nucleic acid ligand-based sandwich assays, designed on two different types of beads that can be readily analysed in flow cytometry, allow multiplexed analysis of a mixture of target protein in a single tube. This sequence is the alpha-thrombin specific DT-3'-LNK-aptamer, a derivative of the dr-aptamer (AAS16530) the capture molecule used to detect alpha-thrombin in a sample using the method described in the invention.	CC
XX	Sequence 43 BP; 6 A; 6 C; 16 G; 15 T; 0 other;	CC
XX	Query Match 100.0%; Score 29; DB 24; Length 43; Best Local Similarity 100.0%; Pred. No. 0.014; Matches 29; Conservative 0; Mismatches 0; Indels 0; Gaps 0;	CC
QY	1 agtcggcggtggggcgggttgggtact 29	CC
Db	5 agtcccggtggggcgggttgggtact 33	CC
RESULT 13		CC
AAT00205		CC
ID AAT00205 standard; DNA; 30 BP.		CC
XX		CC
AC AAT00205;		CC
14-AUG-1996 (first entry)		XX
DE Thrombin DNA ligand, clone #6.		XX
XX		XX
KW family 1; family 2; ligand; thrombin;		XX
KW systematic evolution of ligands by exponential enrichment; SELEX;		XX
KW heparin; selection; region of homology; inhibitor; ss.		XX
OS Synthetic.		XX
PN WO9521853-A1.		XX
XX		XX
PD 17-AUG-1995.		XX
XX		XX
PF 06-FEB-1995; 95WO-US01458.		XX
XX		XX
PR 28-MAR-1994; 94US-0219012.		XX
PR 10-FEB-1994; 94US-0195005.		XX
PR 11-JUN-1990; 90US-0536428.		XX
PR 10-JUN-1991; 91US-0714131.		XX
PR 22-APR-1993; 93US-0061651.		XX
PA (NEXS-) NEXSTAR PHARM INC.		XX
PA Gold L, Janjic N, Tasset D;		XX
PI		XX
XX	Identification of ligands to basic fibroblast growth factor and thrombin - which can be modified for increased in vivo stability	XX
PT		XX
PT	Novel nucleic acid ligands to basic fibroblast growth factor that are useful as inhibitors of basic fibroblast growth factors and 2'-amino modified RNA ligands, exhibit increased in vivo stability	XX
PS	Example 19; Column 57-58; 153pp; English.	PS
XX	WPI; 1995-293073/38.	XX
DR		XX
XX	The sequences given in AAT00202-25 and AAT00227-57 represent two groups of ligands to thrombin. These sequences were isolated using the single stranded DNA molecules given in AAT00201 and AAT00226 which comprise a 3'ON and a 6'ON variable region, respectively. These ligands were isolated using systematic evolution of ligands by exponential enrichment (SELEX). The selection was conducted in a buffer solution at 37 deg. C. After 12 rounds of selection, no additional improvement in binding was seen. By studying regions of homology between the isolated ligands, a truncated ligand of 38 nucleotides (see AAT00810-04) was identified which retains high affinity binding and inhibits clotting. These ligands are inhibitors of thrombin and are therefore useful in treating thrombin mediated conditions and in studying the structure and binding of thrombin.	XX
XX	Sequence 30 BP; 5 A; 5 C; 14 G; 6 T; 0 other;	XX
XX	Query Match 68.3%; Score 19.8; DB 16; Length 30; Best Local Similarity 91.3%; Pred. No. 61; Matches 21; Conservative 0; Mismatches 2; Indels 0; Gaps 0;	XX
QY	4 ccgtgggtggggcgggttgggtgt 26	XX
Db	5 ccgtgggtggggcgggttgggtgt 27	XX
RESULT 14		XX
AAE0757		XX
ID AAE0757 standard; DNA; 30 BP.		XX
XX		XX
AC AAE0757;		XX
XX		XX
DT 20-APR-2001 (first entry)		XX
XX		XX
DE Thrombin high affinity ligand #4.		XX
XX		XX
KW Ligand; basic fibroblast growth factor; bFGF; gene therapy; vascular; atherosclerosis; angioplasty; stability; ss.		XX
KW		XX
XX		XX
OS Unidentified.		XX
XX		XX
PN US617557-B1.		XX
XX		XX
PD 23-JAN-2001.		XX
XX		XX
PF 05-AUG-1996; 96US-0687421.		XX
XX		XX
PR 11-JUN-1990; 90US-0336428.		XX
PR 10-JUN-1991; 91US-0714131.		XX
PR 06-NOV-1992; 92US-073333.		XX
PR 10-FEB-1994; 94US-0195005.		XX
PR 28-MAR-1994; 94US-0219012.		XX
XX		XX
PA (NEXS-) NEXSTAR PHARM INC.		XX
XX		XX
PI Janjic N, Gold L, Tasset D;		XX
XX		XX
DR WPI; 2001-158583/16.		XX
XX		XX
PT Novel nucleic acid ligands to basic fibroblast growth factor that are useful as inhibitors of basic fibroblast growth factors and 2'-amino modified RNA ligands, exhibit increased in vivo stability		XX
PS Example 19; Column 57-58; 153pp; English.		PS
XX		XX

CC The present invention relates to a purified and isolated non-naturally occurring DNA ligands to basic fibroblast growth factor (bFGF).
 CC The ligands are useful as part of gene therapy treatments and
 CC for diagnosing pathogenesis of vascular diseases including
 CC initiation and progression of atherosclerosis, acute coronary
 CC syndromes, vein graft disease and restenosis following coronary
 CC angioplasty. The ligands have improved stability in vivo.
 XX Sequence 30 BP; 5 A; 5 C; 14 G; 6 T; 0 other;

Query Match Best Local Similarity 68.3%; Score 19.8; DB 22; Length 30;
 Matches 21; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy 4 ccgtcgatggcggatggatgggtg 26

Db 5 ccgtcgatggcggatggatgggtg 27

SQ Sequence 30 BP; 4 A; 3 C; 14 G; 9 T; 0 other;
 Query Match Best Local Similarity 66.9%; Score 19.4; DB 16; Length 30;
 Matches 20; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
 Qy 5 cgtcgatggcggatggatgggt 25

Db 9 cgtcgatggcggatggatgggt 29

Search completed: June 6, 2002, 16:08:48
 Job time: 2149 sec

Query Match Best Local Similarity 91.3%; Score 19.8; DB 22; Length 30;
 Matches 21; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy 4 ccgtcgatggcggatggatgggtg 26

Db 5 ccgtcgatggcggatggatgggtg 27

RESULT 15

00218 AAT00218 standard; DNA; 30 BP.

XX AC AAT00218;

XX DT 14-AUG-1996 (first entry)

XX DE Thrombin DNA ligand, clone #26.

XX KW Family 1; ligand; thrombin; systematic evolution of ligands by exponential enrichment; SELEX; heparin; selection; region of homology; inhibitor; ss.

XX OS Synthetic.

XX PN W09521853-A1.

XX PD 17-AUG-1995.

XX PP 06-FEB-1995; 95WO-US01458.

XX PR 28-MAR-1994; 94US-0219012.

PR 10-FEB-1994; 94US-0195005.

PR 11-JUN-1990; 90US-0536428.

PR 10-JUN-1991; 91US-0714131.

PR 22-APR-1993; 93US-0061691.

XX (NEXS-) NEXSTAR PHARM INC.
 XX Gold L, Janjic N, Tasset D;
 XX DR WPI; 1995-293073/38.
 XX PT Identification of ligands to basic fibroblast growth factor and
 PT thrombin - which can be modified for increased in vivo stability
 XX PS Claim 39; Page 95; 236pp; English.

XX The sequences given in AAT00202-25 and AAT00227-57 represent two groups
 CC of ligands to thrombin. These sequences were isolated using the single
 CC stranded DNA molecules given in AAT00201 and AAT00226 which comprise a
 CC 30N and a 60N variable region, respectively. These ligands were
 CC isolated using systematic evolution of ligands by exponential enrichment
 CC (SELEX). The selection was conducted in a buffer solution at 37 deg. C.
 CC After 12 rounds of selection, no additional improvement in binding was
 CC seen. By studying regions of homology between the isolated ligands, a
 CC truncated ligand of 38 nucleotides (see AAQ98403-04) was identified, which
 CC retains high affinity binding and inhibits clotting. These ligands are
 CC inhibitors of thrombin and are therefore useful in treating thrombin
 CC mediated conditions and in studying the structure and binding of
 CC thrombin.

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OM nucleic - nucleic search, using sw model

Run on: June 6, 2002, 16:04:32 ; search time 51.8 Seconds
(without alignments)
137.517 Million cell updates/sec

Title: US-09-599-220-2
Perfect score: 29
Sequence: 1 agtcccggtggcagggtgggtgact 29
Scoring table: IDENTITY-NUC
Gapop 10.0 , Gapext 1.0

Searched: 389533 seqs, 122816752 residues

Total number of hits satisfying chosen parameters: 543772

Minimum DB seq length: 0
Maximum DB seq length: 50

Post-processing: Minimum Match 0%, Maximum Match 100%, Listing first 45 summaries

Database : Issued_Patents_NA:*

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3: /cgn2_6/ptodata/2/ina/6A_COMB.seq:/*
4: /cgn2_6/ptodata/2/ina/6B_COMB.seq:/*
5: /cgn2_6/ptodata/2/ina/PCNUS_COMB.seq:/*
6: /cgn2_6/ptodata/2/ina/backfilesel.seq:/*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match Length	DB ID	Description
1	29	100.0	36	US-08-424-465-3
2	29	100.0	38	US-08-219-012-91
3	29	100.0	38	US-08-316-321-2
4	29	100.0	38	US-08-316-321-2
5	29	100.0	38	US-08-424-465-6
6	29	100.0	38	US-08-687-421-279
7	29	100.0	39	PCT-US95-09237-2
8	29	100.0	39	US-08-479-783-90
9	29	100.0	39	US-08-479-725-90
10	29	100.0	39	US-08-618-691-88
11	29	100.0	39	US-08-973-124-177
12	29	100.0	39	US-08-991-743C-88
13	29	100.0	39	PCT-US96-08014-177
14	29	100.0	40	US-08-434-465-9
15	29	100.0	40	US-08-434-465-13
16	29	100.0	41	US-08-434-465-12
17	29	100.0	41	US-08-434-465-15
18	29	100.0	41	US-08-434-465-15
19	29	100.0	42	US-08-434-465-8
20	29	100.0	42	US-08-434-465-8
21	19	68.3	30	US-08-219-012-31
22	19	68.3	30	US-08-687-421-219
23	19	66.9	30	US-08-213-012-44
24	19	66.9	30	US-08-687-421-232
25	19	65.5	30	US-08-219-012-47
26	19	65.5	30	US-08-687-421-235
27	18	64.8	30	US-08-219-012-29

ALIGNMENTS

RESULT 1
US-08-434-465-3
; Sequence 3, Application US/08434465
; Patent No. 601020
; GENERAL INFORMATION:
; APPLICANT: LARRY GOLD, PAUL G. SCHMIDT, NEBOJSA JANJIC
; TITLE OF INVENTION: NUCLEIC ACID LIGAND COMPLEXES
; NUMBER OF SEQUENCES: 15
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Swanson and Bratschun, L.L.C.
; STREET: 8400 East Prentice Avenue, Suite #200
; CITY: Denver
; STATE: Colorado
; COUNTRY: USA
; ZIP: 80111
; COMPUTER READABLE FORM:
; MEDIUM TYPE: DISKETTE, 3.50 inch, 1.44 Mb storage
; COMPUTER: IBM compatible
; OPERATING SYSTEM: MS-DOS
; SOFTWARE: WORDPERFECT 5.1
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/434,465
; FILING DATE: 4-MAY-1995
; CLASSIFICATION: 514
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 07/714,131
; FILING DATE: 10-JUNE-1991
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 07/536,428
; FILING DATE: 11-JUNE-1990
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 08/234,997
; FILING DATE: 28-April-1994
; ATTORNEY/AGENT INFORMATION:
; NAME: Diana H. McCleard
; REGISTRATION NUMBER: 33,960
; REFERENCE/DOCKET NUMBER: NEX29
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (303) 793-3333
; TELEFAX: (303) 793-3433
; INFORMATION FOR SEQ ID NO: 3:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 36
; REFERENCE/DOCKET NUMBER: NEX29
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (303) 793-3333
; SEQUENCE 21, App
; SEQUENCE 44, App
; SEQUENCE 235, App
; SEQUENCE 29, App
; SEQUENCE 3, App
; SEQUENCE 223, App
; SEQUENCE 43, App
; SEQUENCE 231, App
; SEQUENCE 32, App
; SEQUENCE 240, App
; SEQUENCE 7, App
; SEQUENCE 4, App
; SEQUENCE 64, App
; SEQUENCE 4, App
; SEQUENCE 78, App

SEQUENCE TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: DNA
US-08-434-465-3

Query Match 100.0%; Score 29; DB 3; Length 36;
 Best Local Similarity 100.0%; Pred. No. 0.002; Mismatches 0; Indels 0; Gaps 0; Matches 29; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

RESULT 2
 US-08-219-012-91
 Sequence 91, Application US/08219012
 GENERAL INFORMATION:
 PATENT NO. 5543293
 APPLICANT: Larry Gold
 ADDRESS: Beaton & Swanson, P.C.
 STREET: 4582 South Ulster Street, Parkway, Suite # 403
 CITY: Denver
 STATE: Colorado
 COUNTRY: USA
 ZIP: 80237
 COMPUTER READABLE FORM:
 MEDIUM TYPE: Diskette, 5.25 inch, 360 Kb storage
 COMPUTER: IBM compatible
 OPERATING SYSTEM: MS-DOS
 SOFTWARE: WordPerfect 5.1
 CURRENT APPLICATION DATA:
 APPLICATION NUMBER: US/08/219,012
 FILING DATE:
 CLASSIFICATION: 435
 PRIOR APPLICATION DATA: none
 ATTORNEY/AGENT INFORMATION:
 NAME: Barry J. Swanson
 REGISTRATION NUMBER: 33,215
 REFERENCE/DOCKET NUMBER:
 TELECOMMUNICATION INFORMATION:
 TELEPHONE: (303) 850-9900
 TELEFAX: (303) 850-9401
 INFORMATION FOR SEQ ID NO: 91:
 SEQUENCE CHARACTERISTICS:
 LENGTH: 38 base pairs
 TYPE: nucleic acid
 STRANDEDNESS: single
 TOPOLOGY: linear
 08-219-012-91

Query Match 100.0%; Score 29; DB 1; Length 38;
 Best Local Similarity 100.0%; Pred. No. 0.002; Mismatches 0; Indels 0; Gaps 0; Matches 29; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

RESULT 3
 US-08-376-329-2
 Sequence 91, Application US/08219012
 GENERAL INFORMATION:
 PATENT NO. 5543293
 APPLICANT: Larry Gold
 ADDRESS: Beaton & Swanson, P.C.
 STREET: 4582 South Ulster Street, Parkway, Suite # 403
 CITY: Denver
 STATE: Colorado
 COUNTRY: USA
 ZIP: 80237
 COMPUTER READABLE FORM:
 MEDIUM TYPE: Floppy disk
 COMPUTER: IBM PC compatible
 OPERATING SYSTEM: PC-DOS/MS-DOS
 SOFTWARE: PatentIn Release #1.0, version #1.25
 CURRENT APPLICATION DATA:
 APPLICATION NUMBER: US/08/376,329
 FILING DATE:
 CLASSIFICATION: 435
 ATTORNEY/AGENT INFORMATION:
 NAME: Highet, David W
 REGISTRATION NUMBER: 30,265
 REFERENCE/DOCKET NUMBER: P-3126
 TELECOMMUNICATION INFORMATION:
 TELEPHONE: 201 847 5317
 TELEFAX: 201 848 9228
 INFORMATION FOR SEQ ID NO: 2:
 SEQUENCE CHARACTERISTICS:
 LENGTH: 38 base pairs
 TYPE: nucleic acid
 STRANDEDNESS: single
 TOPOLOGY: linear
 MOLECULE TYPE: DNA (genomic)
 ; US-08-376-329-2

Query Match 100.0%; Score 29; DB 1; Length 38;
 Best Local Similarity 100.0%; Pred. No. 0.002; Mismatches 0; Indels 0; Gaps 0; Matches 29; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

RESULT 4
 US-08-276-271-2
 Sequence 2, Application US/08276271
 GENERAL INFORMATION:
 PATENT NO. 5650275
 APPLICANT: Pitner, James B
 APPLICANT: Malinowski, Douglas P
 APPLICANT: Vonk, Glenn P
 APPLICANT: Gold, Larry
 TITLE OF INVENTION: Spectroscopically Detectable Nucleic Acid Ligands
 NUMBER OF SEQUENCES: 4
 CORRESPONDENCE ADDRESS:
 ADDRESSEE: Richard J. Roderick, Becton Dickinson and Company
 STREET: 1 Becton Drive
 CITY: Franklin Lakes
 STATE: NJ
 COUNTRY: USA
 ZIP: 07417-1880
 COMPUTER READABLE FORM:
 MEDIUM TYPE: Floppy disk
 COMPUTER: IBM PC compatible
 OPERATING SYSTEM: PC-DOS/MS-DOS
 SOFTWARE: PatentIn Release #1.0, version #1.25
 CURRENT APPLICATION DATA:
 APPLICATION NUMBER: US/08/276,271
 FILING DATE:
 CLASSIFICATION: 435
 ATTORNEY/AGENT INFORMATION:

NAME: Hight, David W
 REGISTRATION NUMBER: 30,265
 REFERENCE/DOCKET NUMBER: P-3126
 TELECOMMUNICATION INFORMATION:
 TELEPHONE: 201 847 5317
 TELEFAX: 201 848 9228
 INFORMATION FOR SEQ ID NO: 2:
 SEQUENCE CHARACTERISTICS:
 LENGTH: 38 base pairs
 TYPE: nucleic acid
 STRANDEDNESS: single
 TOPOLOGY: linear
 MOLECULE TYPE: DNA (genomic)
 US-08-276-271-2

Query Match, Best Local Similarity 100.0%; Score 29; DB 1; Length 38; Matches 29; Conservative 0; Mismatches 0; Indels 0; Gaps 0; STRANDEDNESS: single
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 2 AGTCCGGTGGCAGGTGGGTGACT 30

RESULT 5
 US-08-134-465-6
 Sequence 6, Application US/08434465
 PATENT NO. 6011020
 GENERAL INFORMATION:
 APPLICANT: LARRY GOLD, PAUL G. SCHMIDT, NEBOJSA JANJIC
 TITLE OF INVENTION: NUCLEIC ACID LIGAND COMPLEXES
 NUMBER OF SEQUENCES: 15
 CORRESPONDENCE ADDRESS:
 ADDRESSEE: Swanson and Bratschun, L.L.C.
 STREET: 8400 East Prentice Avenue, Suite #200
 CITY: Denver
 STATE: Colorado
 COUNTRY: USA
 ZIP: 80111
 COMPUTER READABLE FORM:
 COMPUTER: IBM compatible
 OPERATING SYSTEM: MS-DOS
 SOFTWARE: WordPerfect 5.1
 CURRENT APPLICATION DATA:
 APPLICATION NUMBER: US/08/434,465
 FILING DATE: 4-MAY-1995
 CLASSIFICATION: 514
 PRIORITY APPLICATION DATA:
 APPLICATION NUMBER: 07/714,131
 FILING DATE: 10-JUNE-1991
 PRIORITY APPLICATION DATA:
 APPLICATION NUMBER: 07/536,428
 FILING DATE: 11-JUNE-1990
 PRIORITY APPLICATION DATA:
 APPLICATION NUMBER: 08/234,997
 FILING DATE: 28-APRIL-1994
 ATTORNEY/AGENT INFORMATION:
 NAME: Diane H. McClearn
 REGISTRATION NUMBER: 33,950
 REFERENCE/DOCKET NUMBER: NEX29
 TELECOMMUNICATION INFORMATION:
 TELEPHONE: (303) 793-3333
 TELEFAX: (303) 793-3433
 INFORMATION FOR SEQ ID NO: 6:
 SEQUENCE CHARACTERISTICS:
 LENGTH: 38
 TYPE: nucleic acid
 STRANDEDNESS: single
 TOPOLOGY: linear
 MOLECULE TYPE: DNA
 FEATURE: OTHER INFORMATION: Nucleotides 37 and 38 are bound by a phosphorothioate bond

RESULT 6
 US-08-687-421-279
 Sequence 279, Application US/08687421
 PATENT NO. 617757
 GENERAL INFORMATION:
 APPLICANT: Gold, Larry
 APPLICANT: Janjic, Nebojsa
 APPLICANT: Tasset, Diane
 TITLE OF INVENTION: HIGH-AFFINITY LIGANDS OF BASIC FIBROBLAST GROWTH FACTOR AND TITLE OF INVENTION: THROMBIN
 NUMBER OF SEQUENCES: 445
 CORRESPONDENCE ADDRESS:
 ADDRESSEE: Swanson & Bratschun, L.L.C.
 STREET: 8400 E. Prentice Avenue, Suite 200
 CITY: Englewood
 STATE: Colorado
 COUNTRY: USA
 ZIP: 80111
 COMPUTER READABLE FORM:
 COMPUTER: IBM compatible, 3.5 inch, 1.44 MB storage
 OPERATING SYSTEM: MS-DOS
 SOFTWARE: WordPerfect 6.0
 CURRENT APPLICATION DATA:
 APPLICATION NUMBER: US/08/687,421
 FILING DATE: 08-MAY-1996
 CLASSIFICATION: 435
 PRIORITY APPLICATION DATA:
 APPLICATION NUMBER: 08/195,005
 FILING DATE: 10-FEBRUARY-1994
 PRIORITY APPLICATION DATA:
 APPLICATION NUMBER:
 FILING DATE: 22-APRIL-1993
 PRIORITY APPLICATION DATA:
 APPLICATION NUMBER: 08/219,012
 FILING DATE: 28-MARCH-1994
 PRIORITY APPLICATION DATA:
 APPLICATION NUMBER: 07/973,333
 FILING DATE: 11-NOVEMBER-1992
 PRIORITY APPLICATION DATA:
 APPLICATION NUMBER: 07/714,131
 FILING DATE: 10-JUNE-1991
 PRIORITY APPLICATION DATA:
 APPLICATION NUMBER: 07/536,428
 FILING DATE: 11-JUNE-1990
 ATTORNEY/AGENT INFORMATION:
 NAME: Barry J. Swanson
 REGISTRATION NUMBER: 33,215
 REFERENCE/DOCKET NUMBER: NEX07/PCT
 TELECOMMUNICATION INFORMATION:
 TELEPHONE: (303) 793-3333
 TELEFAX: (303) 793-3433
 INFORMATION FOR SEQ ID NO: 279:
 SEQUENCE CHARACTERISTICS:
 LENGTH: 38 base pairs
 TYPE: nucleic acid
 STRANDEDNESS: single
 TOPOLOGY: linear

US-08-687-421-279

ADDRESSEE: Swanson and Bratschun, L.L.C.
 STREET: 8400 East Prentice Avenue, Suite #200
 CITY: Denver
 STATE: Colorado
 COUNTRY: USA
 ZIP: 80111
 COMPUTER READABLE FORM:
 MEDIUM TYPE: Diskette, 3.5 inch, 1.4 Mb storage
 COMPUTER: IBM compatible
 OPERATING SYSTEM: MS-DOS
 SOFTWARE: Wordperfect 5.1
 CURRENT APPLICATION DATA:
 APPLICATION NUMBER: US/08/479,725
 FILING DATE: 7-JUNE-1995
 CLASSIFICATION: 536
 PRIORITY APPLICATION DATA:
 APPLICATION NUMBER: 07/114,131
 FILING DATE: 10-JUNE-1991
 PRIORITY APPLICATION DATA:
 APPLICATION NUMBER: 07/931,473
 FILING DATE: 17-AUGUST-1992
 PRIORITY APPLICATION DATA:
 APPLICATION NUMBER: 07/964,624
 FILING DATE: 21-OCTOBER-1992
 PRIORITY APPLICATION DATA:
 APPLICATION NUMBER: 08/117,991
 FILING DATE: 8-SEPTEMBER-1993
 PRIORITY APPLICATION DATA:
 APPLICATION NUMBER: 07/536,428
 FILING DATE: 11-JUNE-1990
 ATTORNEY/AGENT INFORMATION:
 NAME: Diane H. McClearn
 REGISTRATION NUMBER: 33,960
 REFERENCE/DOCKET NUMBER: NECX42-1
 TELECOMMUNICATION INFORMATION:
 TELEPHONE: (303) 793-3333
 TELEFAX: (303) 793-3433
 INFORMATION FOR SEQ ID NO: 90:
 SEQUENCE CHARACTERISTICS:
 LENGTH: 39 base pairs
 TYPE: nucleic acid
 STRANDEDNESS: single
 TOPOLOGY: linear
 MOLECULE TYPE: DNA
 FEATURE:
 OTHER INFORMATION: Nucleotide 39 is an inverted
 sequence 88, Application US/08618693
 LT 10
 LT 618-693-88
 LT Local Similarity 100.0%; Score 29; DB 1; Lu
 tient No. 5123594
 GENERAL INFORMATION:
 APPLICANT: NEBOUSA JANJIC
 APPLICANT: LARRY GOLD
 TITLE OF INVENTION: HIGH AFFINITY PDGF NUCLEIC
 TITLE OF INVENTION: ACID LIGANDS
 NUMBER OF SEQUENCES: 96
 CORRESPONDENCE ADDRESS:
 ADDRESSEE: Swanson and Bratschun, L.L.C.
 STREET: 8400 East Prentice Avenue, Suite #200
 CITY: Denver

COMPUTER READABLE FORM:
MEDIUM TYPE: *Diskette*, 3.5 inch, 1.4 Mb storage
COMPUTER: *IBM* compatible
OPERATING SYSTEM: *MS-DOS*
SOFTWARE: *Wordperfect* 6.0
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US 08/618,693
FILING DATE: 20 MARCH 1996
CLASSIFICATION: 536
PRIORITY APPLICATION DATA:
APPLICATION NUMBER: 08/479,783
FILING DATE: 7-JUNE-1995
ATTORNEY/AGENT INFORMATION:
NAME: *Barry J. Swanson*
REGISTRATION NUMBER: 33,215
REFERENCE/DOCKET NUMBER: *NEW42/CIP*
TELECOMMUNICATION INFORMATION:
TELEPHONE: (303) 793-3333
TELEFAX: (303) 793-3433
INFORMATION FOR SEQ ID NO: 88:
SEQUENCE CHARACTERISTICS:
LENGTH: 39 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: DNA
FEATURE:
OTHER INFORMATION: Nucleotide 39 is an inverted orientation T (3 - 3 linked)
OTHER INFORMATION: US-08-618-693-88

REGISTRATION NUMBER: 33,960
REFERENCE DOCKET NUMBER: NEX29
TELECOMMUNICATION INFORMATION:
TELEPHONE: (303) 793-3333
TELEFAX: (303) 793-3433
INFORMATION FOR SEQ ID NO: 13:
SEQUENCE CHARACTERISTICS:
LENGTH: 40
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: DNA
FEATURE:
OTHER INFORMATION: N at position 1 is a 20,000 MW
OTHER INFORMATION: PEG
FEATURE:
OTHER INFORMATION: N at position 38 is a dT amino
OTHER INFORMATION: phosphoramidite
FEATURE:
OTHER INFORMATION: Nucleotide 39 is an inverted
OTHER INFORMATION: orientation (3',3' linkage) phosphoramidite
85-08-434-465-13

Query Match 100.0%; Score 29; DB 3; Length 40;
Best Local Similarity 100.0%; Pred. No. 0.002; Mismatches 0; Indels 0; Gaps 0;
Matches 29; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 agtcgcgtggtagggcagggtgggtgact 29
Db 1 ||||||| ||||||| ||||||| ||||||| ||||||| |||||
3 AGTCGGTAGGGCAGGGTGGGTGACT 31

Search completed: June 6, 2002, 16:04:32
Job time: 1893 sec

Gencore version 4.5
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On nucleic - nucleic search, using sw model

Run on: June 6, 2002, 15:32:59 ; search time 1796.86 Seconds

(Without alignments)
 174.693 Million cell updates/sec

Title: US-09-599-220-1

Perfect score: 15

Sequence: 1 ggttagtggtgg 15

Scoring table: IDENTITY_NUC

Gabop 10.0 , Gapext 1.0

Searched: 1797656 seqs, 10463268293 residues

Total number of hits satisfying chosen parameters: 708260

Minimum DB seq length: 0

Maximum DB seq length: 50

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

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1: qb_bp:*

2: qb_hsq:*

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4: qb_om:*

5: qb_ov:*

6: qb_pat:*

7: qb_phn:*

8: qb_pl:*

9: qb_pr:*

10: qb_1o:*

11: qb_sts:*

12: qb_sy:*

13: qb_un:*

14: qb_v1:*

15: em_ba:*

16: em_fun:*

17: em_hum:*

18: em_in:*

19: em_mu:*

20: em_on:*

21: em_or:*

22: em_ov:*

23: em_pat:*

24: em_ph:*

25: em_pl:*

26: em_ro:*

27: em_sts:*

28: em_un:*

29: em_v1:*

30: em_hg_hum:*

31: em_hg_inv:*

32: em_hg_other:*

33: em_higo_inv:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

ALIGNMENTS

RESULT

1

AR009266

LOCUS AR009266

DEFINITION Sequence 29 from patent US 5756291.

ACCESSION AR009266

VERSION AR009266.1

KEYWORDS

Unknown.

SOURCE ORGANISM

Unclassified.

REFERENCE 1 (bases 1 to 15)

AUTHORS Griffin,L., Albrecht,G., Latham,J., Leung,L., Vermaas,E. and

TITLE Aptamers specific for biomolecules and methods of making

JOURNAL Patent: US 5756291-A 29 26 MAY 1998;

FEATURES Location/Qualifiers

source

1..15

/organism="Unknown"

SUMMARIES

Result No.	Score	Query Match Length	DB ID	Description
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Query	Match	Length	DB	ID
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Query	Match	Length	DB	ID
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Best Local Similarity 100.0%; Pred. No. 3e+03; Mismatches 0; Indels 0; Gaps 0; Source Unknown.

DEFINITION Sequence 36 from patent US 5756291.

ACCESSION AR009273

VERSION AR09273.1 GI:3968078

KEYWORDS

ORGANISM Unknown.

Unclassified.

1 (bases 1 to 15).

REFERENCE

AUTHORS Griffin,L., Albrecht,G., Latham,J., Leung,L., Vermaas,E. and Toole,J.J.

TITLE Aptamers specific for biomolecules and methods of making

JOURNAL Patent: US 5756291-A 31 26-MAY-1998;

FEATURES Location/Qualifiers 1..15

KEYWORDS /organism="unknown"

SOURCE 1.15

BASE COUNT 0 a 0 c 9 g 6 t

ORIGIN

RESULT 2

Query Match 100.0%; Score 15; DB 6; Length 15;

LOCUS 15 bp DNA linear PAT 04-DEC-1998

DEFINITION Sequence 31 from patent US 5756291.

ACCESSION AR009268

VERSION AR009268.1 GI:3968073

KEYWORDS

ORGANISM Unknown.

Unclassified.

1 (bases 1 to 15).

REFERENCE

AUTHORS Griffin,L., Albrecht,G., Latham,J., Leung,L., Vermaas,E. and Toole,J.J.

TITLE Aptamers specific for biomolecules and methods of making

JOURNAL Patent: US 5756291-A 36 26-MAY-1998;

FEATURES Location/Qualifiers 1..15

KEYWORDS /organism="unknown"

SOURCE 1.15

BASE COUNT 0 a 0 c 9 g 6 t

ORIGIN

RESULT 3

Query Match 100.0%; Score 15; DB 6; Length 15;

LOCUS 15 bp DNA linear PAT 04-DEC-1998

DEFINITION Sequence 32 from patent US 5756291.

ACCESSION AR009269

VERSION AR009269.1 GI:3968074

KEYWORDS

SOURCE Unknown.

ORGANISM Unknown.

Unclassified.

1 (bases 1 to 15).

REFERENCE

AUTHORS Griffin,L., Albrecht,G., Latham,J., Leung,L., Vermaas,E. and Toole,J.J.

TITLE Aptamers specific for biomolecules and methods of making

JOURNAL Patent: US 5756291-A 32 26-MAY-1998;

FEATURES Location/Qualifiers 1..15

KEYWORDS /organism="unknown"

SOURCE 1.15

BASE COUNT 0 a 0 c 9 g 6 t

ORIGIN

RESULT 4

Query Match 100.0%; Score 15; DB 6; Length 15;

LOCUS 15 bp DNA linear PAT 04-DEC-1998

DEFINITION Sequence 38 from patent US 5756291.

ACCESSION AR009275

VERSION AR009275.1 GI:3968080

KEYWORDS

ORGANISM Unknown.

Unclassified.

1 (bases 1 to 15).

REFERENCE

AUTHORS Griffin,L., Albrecht,G., Latham,J., Leung,L., Vermaas,E. and Toole,J.J.

TITLE Aptamers specific for biomolecules and methods of making

RESULT 5

Query Match 100.0%; Score 15; DB 6; Length 15;

LOCUS 15 bp DNA linear PAT 04-DEC-1998

DEFINITION Sequence 37 from patent US 5756291.

ACCESSION AR009274

VERSION AR009274.1 GI:3968079

KEYWORDS

ORGANISM Unknown.

Unclassified.

1 (bases 1 to 15).

REFERENCE

AUTHORS Griffin,L., Albrecht,G., Latham,J., Leung,L., Vermaas,E. and Toole,J.J.

TITLE Aptamers specific for biomolecules and methods of making

JOURNAL Patent: US 5756291-A 37 26-MAY-1998;

FEATURES Location/Qualifiers 1..15

KEYWORDS /organism="unknown"

SOURCE 1.15

BASE COUNT 0 a 0 c 9 g 6 t

ORIGIN

RESULT 6

Query Match 100.0%; Score 15; DB 6; Length 15;

LOCUS 15 bp DNA linear PAT 04-DEC-1998

DEFINITION Sequence 38 from patent US 5756291.

ACCESSION AR009275

VERSION AR009275.1 GI:3968080

KEYWORDS

ORGANISM Unknown.

Unclassified.

1 (bases 1 to 15).

REFERENCE

AUTHORS Griffin,L., Albrecht,G., Latham,J., Leung,L., Vermaas,E. and Toole,J.J.

TITLE Aptamers specific for biomolecules and methods of making

RESULT 7

Query Match 100.0%; Score 15; DB 6; Length 15;

LOCUS 15 bp DNA linear PAT 04-DEC-1998

DEFINITION Sequence 38 from patent US 5756291.

ACCESSION AR009275

VERSION AR009275.1 GI:3968080

KEYWORDS

ORGANISM Unknown.

Unclassified.

1 (bases 1 to 15).

REFERENCE

AUTHORS Griffin,L., Albrecht,G., Latham,J., Leung,L., Vermaas,E. and Toole,J.J.

TITLE Aptamers specific for biomolecules and methods of making

JOURNAL Patent: US 556291-A 38 26-MAY-1998;
 FEATURES Location/Qualifiers
 source 1.
 /organism="unknown"
 BASE COUNT 0 a 0 c 9 g 6 t
 ORIGIN

RESULT 9
 QY 1 ggttgggtgtgtgg 15
 LOCUS AR009316 15 bp DNA linear PAT 29-SEP-1998
 DEFINITION Sequence 21 from patent US 5840867.
 ACCESSION AR060777.1 GI:5987227
 VERSION AR060777
 KEYWORDS .
 SOURCE Unknown.
 ORGANISM Unknown.
 UNCLASSIFIED Unclassified.
 REFERENCE 1. (bases 1 to 15)
 AUTHORS Toole,J.J., Griffin,L.C., Bock,L.C. and Latham,J.A.
 TITLE Aptamers specific for biomolecules and methods of making
 JOURNAL Patent: US 5756291-A 79 26-MAY-1998;
 ACCESSION AR009316.1 GI:3968121
 VERSION 1.
 FEATURES Location/Qualifiers 1..15
 source /organism="unknown"
 BASE COUNT 0 a 0 c 9 g 6 t
 ORIGIN

RESULT 7
 QY 1 ggttgggtgtgtgg 15
 LOCUS AR009316 15 bp DNA linear PAT 04-DEC-1998
 DEFINITION Sequence 79 from patent US 5756291.
 ACCESSION AR060777
 VERSION AR009316.1
 KEYWORDS .
 SOURCE Unknown.
 ORGANISM Unknown.
 UNCLASSIFIED Unclassified.
 REFERENCE 1. (bases 1 to 15)
 AUTHORS Griffin,L., Albrecht,G., Latham,J., Leung,L., Vermaas,E. and
 Toole,J.J.
 TITLE Aptamers specific for biomolecules and methods of making
 JOURNAL Patent: US 5756291-A 79 26-MAY-1998;
 ACCESSION AR009316.1
 VERSION 1..15
 FEATURES Location/Qualifiers 1..15
 source /organism="unknown"
 BASE COUNT 0 a 0 c 9 g 6 t
 ORIGIN

RESULT 10
 QY 1 ggttgggtgtgtgg 15
 LOCUS AR060778 15 bp DNA linear PAT 29-SEP-1999
 DEFINITION Sequence 22 from patent US 5840867.
 ACCESSION AR060778
 VERSION AR060778.1 GI:5987228
 KEYWORDS .
 SOURCE Unknown.
 ORGANISM Unknown.
 UNCLASSIFIED Unclassified.
 REFERENCE 1. (bases 1 to 15)
 AUTHORS Toole,J.J., Griffin,L.C., Bock,L.C. and Latham,J.A.
 TITLE Aptamer analogs specific for biomolecules
 JOURNAL Patent: US 5840867-A 22 24-NOV-1998;
 ACCESSION AR060775
 VERSION AR060775.1 GI:5987225
 KEYWORDS .
 SOURCE Unknown.
 ORGANISM Unknown.
 UNCLASSIFIED Unclassified.
 REFERENCE 1. (bases 1 to 15)
 AUTHORS Toole,J.J., Griffin,L.C., Bock,L.C. and Latham,J.A.
 TITLE Aptamer analogs specific for biomolecules
 JOURNAL Patent: US 5840867-A 19 24-NOV-1998;
 ACCESSION AR060775
 VERSION AR060775.1
 FEATURES Location/Qualifiers 1..15
 source /organism="unknown"
 BASE COUNT 0 a 0 c 9 g 6 t
 ORIGIN

RESULT 8
 QY 1 ggttgggtgtgtgg 15
 LOCUS AR060775 15 bp DNA linear PAT 29-SEP-1999
 DEFINITION Sequence 19 from patent US 5840867.
 ACCESSION AR060775
 VERSION AR060775.1 GI:5987225
 KEYWORDS .
 SOURCE Unknown.
 ORGANISM Unknown.
 UNCLASSIFIED Unclassified.
 REFERENCE 1. (bases 1 to 15)
 AUTHORS Toole,J.J., Griffin,L.C., Bock,L.C. and Latham,J.A.
 TITLE Aptamer analogs specific for biomolecules
 JOURNAL Patent: US 5840867-A 19 24-NOV-1998;
 ACCESSION AR060775
 VERSION AR060775.1
 FEATURES Location/Qualifiers 1..15
 source /organism="unknown"
 BASE COUNT 0 a 0 c 9 g 6 t
 ORIGIN

RESULT 11
 QY 1 ggttgggtgtgtgg 15
 LOCUS AR098723 15 bp DNA linear PAT 14-FEB-2001
 DEFINITION Sequence 81 from patent US 6077668.
 ACCESSION AR098723
 VERSION AR098723.1 GI:12808489
 KEYWORDS .
 SOURCE Unknown.

Query Match 100.0%; Score 15; DB 6; Length 15;
 Best Local Similarity 100.0%; Pred. No. 3e+03; 0; Mismatches
 Matches 15; Conservative 0; Indels 0; Gaps 0;

ORGANISM Unknown.
REFERENCE Unclassified.
AUTHORS 1 (bases 1 to 15)
TITLE Highly sensitive multimeric nucleic acid probes
JOURNAL Patent: US 6077668-A 01-20-JUN-2000;
FEATURES 1..15
source /organism="unknown"
BASE COUNT 0 a 0 c 9 g 6 t
ORIGIN

RESULT 12
AR125847
LOCUS AR125847 15 bp DNA linear PAT 16-MAY-2001
DEFINITION Sequence 15 from patent US 6177557.
ACCESSION AR125847
VERSION AR125847.1 GI:14111909
KEYWORDS
ORGANISM Unknown.
REFERENCE 1 (bases 1 to 15)
AUTHORS Janjic,N., Gold,L. and Tasse,D.
TITLE High affinity ligands of basic fibroblast growth factor and
FEATURES thrombin
source /organism="unknown"
BASE COUNT 0 a 0 c 9 g 6 t
ORIGIN

Query Match 100 0%; Score 15; DB 6; Length 15;
Best Local Similarity 100 0%; Pred. No. 3e+03; 0; Mismatches 0; Indels 0; Gaps 0;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 gggtgggtgggtgg 15
Db 1 GGTTGGCTGGTGG 15

RESULT 14
I16587
LOCUS I16587 15 bp DNA linear PAT 03-APR-1996
DEFINITION Sequence 1 from patent US 5476766.
ACCESSION I16587
VERSION I16587.1 GI:1251495
KEYWORDS
ORGANISM Unknown.
REFERENCE 1 (bases 1 to 15)
AUTHORS Gold,L. and Tasse,D.
TITLE Ligands of thrombin
FEATURES source /organism="unknown"
BASE COUNT 0 a 0 c 9 g 6 t
ORIGIN

Query Match 100 0%; Score 15; DB 6; Length 15;
Best Local Similarity 100 0%; Pred. No. 3e+03; 0; Mismatches 0; Indels 0; Gaps 0;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 gggtgggtgggtgg 15
Db 1 GGTTGGCTGGTGG 15

RESULT 15
I24214
LOCUS I24214 15 bp DNA linear PAT 07-OCT-1996
DEFINITION Sequence 1 from patent US 5543293.
ACCESSION I24214
VERSION I24214.1 GI:1604084
KEYWORDS
ORGANISM Unknown.
REFERENCE 1 (bases 1 to 15)
AUTHORS Gold,L. and Tasse,D.
TITLE DNA ligands of thrombin
JOURNAL Patent: US 5543293-A 1 06-AUG-1996;
FEATURES source /organism="unknown"
BASE COUNT 0 a 0 c 9 g 6 t
ORIGIN

Query Match 100 0%; Score 15; DB 6; Length 15;
Best Local Similarity 100 0%; Pred. No. 3e+03; 0; Mismatches 0; Indels 0; Gaps 0;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 gggtgggtgggtgg 15
Db 1 GGTTGGCTGGTGG 15

RESULT 13
AR168827
LOCUS AR168827 15 bp DNA linear PAT 17-DEC-2001
DEFINITION Sequence 15 from patent US 6288042.
ACCESSION AR168827
VERSION AR168827.1 GI:17904949
KEYWORDS
ORGANISM Unknown.
REFERENCE 1 (bases 1 to 15)
AUTHORS Rando,R.F., O'waug,J.O., Hogan,M.E., Wallace,T.L. and Cossom,P.A.
TITLE Anti-viral guanosine-rich teard forming oligonucleotides
JOURNAL Patent: US 6288042-A 53-11-SEP-2001;
FEATURES source /organism="unknown"
BASE COUNT 0 a 0 c 9 g 6 t
ORIGIN

Search completed: June 6, 2002, 16:03:19
Job time: 1820 sec

Fri Jun 7 09:51:14 2002

us-09-599-220-1.rge

GenCore version 4.5
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OM nucleic - nucleic search, using sw model

Run on: June 6, 2002, 15:32:59 ; Search time 234.25 Seconds
 (without alignments)
 109.941 Million cell updates/sec

Title: US-09-599-220-1
 Perfect score: 15
 Sequence: 1 ggttgggtgtggtttg 15

Scoring table: IDENTITY_NUC
 Gapop 10.0 , Gapext 1.0

Searched: 1736436 seqs, 858457221 residues

Total number of hits satisfying chosen parameters: 1905168

Minimum DB seq length: 0
 Maximum DB seq length: 50

Post-processing: Minimum Match 0%
 Maximum Match 100%
 Listing first 45 summaries

Database : N_Geneseq-032802:*

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 4: /SIDS1/gcgatata/geneseq/geneseq/geneseq-emb1/NA1983.DAT: *
 5: /SIDS1/gcgatata/geneseq/geneseq-emb1/NA1984.DAT: *
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 24: /SIDS1/gcgatata/geneseq/geneseq-emb1/NA2002.DAT: *

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

RESULT 1

ID AAQ28472 standard; DNA; 15 BP.
 XX AAQ28472;
 AC
 XX DT 16-FEB-1993 (first entry)
 XX DE Thrombin aptamer.
 XX KW Aptamer; specifically binding oligonucleotides; primer/linker; PCR; cleavage; ss.
 XX KW
 OS Synthetic.
 XX PN WO9214843-A.
 XX PD 03-SEP-1992.
 XX PF 21-FEB-1992; 92WO-US01383.
 XX PR 21-FEB-1991; 91US-0658796.
 PR 21-FEB-1991; 91US-0658849.
 PR 21-FEB-1991; 91US-0659103.
 PR 21-FEB-1991; 91US-0659113.
 PR 21-FEB-1991; 91US-0659114.
 PR 21-FEB-1991; 91US-0659180.
 PR 21-FEB-1991; 91US-0659181.
 PR 14-AUG-1991; 91US-0744870.
 PR 14-AUG-1991; 91US-0745215.
 PR 06-NOV-1991; 91US-0787921.

ALIGNMENTS

PA (GILE-) GILEAD SCI INC.
 XX PR 14-AUG-1991; 91US-0744870.
 PI Bock LC, Griffin LC, Krawczyk S, Latham JA, Toole JJ; PR 14-AUG-1991; 91US-0744215.
 PT Muenchau DD; PR 06-NOV-1991; 91US-0787921.
 XX XX PA (GILE-) GILEAD SCI INC.
 DR WPI; 1992-316194/38.
 XX PR DNA aptamers specifically binding target molecules - useful for retrieving target molecules, delivering drugs or toxins to desired targets and for treating auto-immune diseases
 PT CC disclosure; Page 119; 183PP; English.
 CC The sequences given in AAQ28473-78 are aptamers which are based on the unmodified thrombin aptamer given in AAQ28472. These aptamers bind to thrombin inhibiting its activity, except for the aptamer sequence given in AAQ28475 which was required in very large quantities to inhibit thrombin activity. These aptamers are stable, versatile and highly specific to their intended targets. They can be used to deliver auxiliary substances, e.g. drugs, toxins, radio isotopes etc. to a specific part of the body. The aptamers have a binding region of approx. 10 nucleotide residues.
 CC XX Sequence 15 BP; 0 A; 0 C; 9 G; 6 T; 0 other;
 PR Query Match 100.0%; Score 15; DB 13; Length 15;
 Best Local Similarity 100.0%; Pred. No. 4.4e+02;
 Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 OY DB 1 ggttgggtgggtgg 15
 XX
 RESULT 2
 AAQ28474
 ID AAQ28474 standard; DNA; 15 BP.
 XX AC AAQ28474;
 XX DT 16-FEB-1993 (first entry)
 XX DE Modified thrombin aptamer #2.
 XX KW Aptamer; specifically binding oligonucleotides; primer/linker; PCR;
 KW cleavage; ss.
 XX Synthetic.
 XX
 FH Key Location/qualifiers
 FT misc_difference 13..14
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 FT /*note= "The linkage between T13 and G14 is a
 FT thioate linkage"
 FT misc_difference 14..15
 FT /*tag= b
 FT /*note= "The linkage between G14 and G15 is a
 FT thioate linkage"
 PN W09214843-A.
 XX 03-SEP-1992.
 XX 21-FEB-1992; 92WO-US01383.
 XX PR 21-FEB-1991; 91US-0558795.
 PR 21-FEB-1991; 91US-0659809.
 PR 21-FEB-1991; 91US-0659813.
 PR 21-FEB-1991; 91US-0659113.
 PR 21-FEB-1991; 91US-0659114.
 PR 21-FEB-1991; 91US-0659980.
 PR 21-FEB-1991; 91US-0659981.
 PR
 XX PR DNA aptamers specifically binding target molecules - useful for retrieving target molecules, delivering drugs or toxins to desired targets and for treating auto-immune diseases
 PT CC disclosure; Page 119; 183PP; English.
 CC The sequences given in AAQ28473-78 are aptamers which are based on the unmodified thrombin aptamer given in AAQ28472. These aptamers bind to thrombin inhibiting its activity, except for the aptamer sequence given in AAQ28475 which was required in very large quantities to inhibit thrombin activity. These aptamers are stable, versatile and highly specific to their intended targets. They can be used to deliver auxiliary substances, e.g. drugs, toxins, radio isotopes etc. to a specific part of the body. The aptamers have a binding region of approx. 10 nucleotide residues.
 CC XX Sequence 15 BP; 0 A; 0 C; 9 G; 6 T; 0 other;
 CC XX PR Query Match 100.0%; Score 15; DB 13; Length 15;
 Best Local Similarity 100.0%; Pred. No. 4.4e+02;
 Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 OY DB 1 ggttgggtgggtgg 15
 XX
 RESULT 3
 AAQ28477
 ID AAQ28477 standard; DNA; 15 BP.
 XX AC AAQ28477;
 XX DT 16-FEB-1993 (first entry)
 XX DE Modified thrombin aptamer #5.
 XX KW Aptamer; specifically binding oligonucleotides; primer/linker; PCR;
 KW cleavage; ss.
 XX OS Synthetic.
 XX
 FH Key Location/qualifiers
 FT misc_difference 3
 FT /*tag= a
 FT /*label= 5-(1-pentyryl)-2'-deoxyuridine
 FT modified_base 12
 FT /*tag= b
 FT /*label= 5-(1-pentyryl)-2'-deoxyuridine
 FT
 XX PN W09214843-A.
 XX PR 21-FEB-1992; 92WO-US01383.
 XX PD 03-SEP-1992.
 XX PR 21-FEB-1992; 92WO-US01383.
 XX PR 21-FEB-1991; 91US-0658796.
 PR 21-FEB-1991; 91US-065849.
 PR 21-FEB-1991; 91US-065903.
 PR 21-FEB-1991; 91US-0659113.
 PR 21-FEB-1991; 91US-0659114.

PR 21-FEB-1991; 91US-0559980.
 PR 21-FEB-1991; 91US-0559981.
 PR 14-AUG-1991; 91US-0744870.
 PR 14-AUG-1991; 91US-0745215.
 PR 06-NOV-1991; 91US-0787921.
 XX PA (GILE-) GILEAD SCI INC.
 XX PI Bock LC, Griffin LC, Krawczyk S, Latham JA, Toole JJ;
 PI Muenchau DD;
 XX DR WPI; 1992-316194/38.
 XX PS DNA aptamers specifically binding target molecules - useful for
 PT retrieving target molecules, delivering drugs or toxins to
 PT desired targets and for treating auto-immune diseases
 XX Disclosure; Page 119; 183pp; English.
 XX CC The sequences given in AAQ28473-78 are aptamers which are based on the
 CC unmodified thrombin aptamer given in AAQ28472. These aptamers bind
 CC to thrombin inhibiting its activity, except for the aptamer sequence
 CC given in AAQ28475 which was required in very large quantities to inhibit
 CC thrombin activity. These aptamers are stable, versatile and highly
 CC specific to their intended targets. They can be used to deliver
 CC auxiliary substances, e.g. drugs, toxins, radio isotopes etc. to a
 CC specific part of the body. The aptamers have a binding region of
 CC approx. 10 nucleotide residues.
 XX SQ Sequence 15 BP; 0 A; 0 C; 9 G; 4 T; 0 other;
 Query Match 100.0%; Score 15; DB 13; Length 15;
 Best Local Similarity 86.7%; Pred. No. 4.4e+02;
 Matches 13; Conservative 2; Mismatches 0; Indels 0; Gaps 0;
 QY 1 ggttgggtgggtgg 15
 Db 1 gggttgggtgggtgg 15
 RESULT 4
 AAQ28478
 ID AAQ28478 standard; DNA; 15 BP.
 XX
 AC AAQ28478;
 XX DT 16-FEB-1993 (first entry)
 XX KW Modified thrombin aptamer #6.
 KW Aptamer; specifically binding oligonucleotides; primer/linker; PCR;
 KW cleavage; ss.
 XX OS Synthetic.
 XX Key Location/Qualifiers
 FT modified_base 13
 FT /*tag= a
 FT /label= 5-(1-pentynyl)uracil
 PN W09214843-A.
 XX PD 03-SEP-1992.
 XX PF 21-FEB-1992; 92WO-US01383.
 XX PR 21-FEB-1991; 91US-0658849.
 PR 21-FEB-1991; 91US-0659103.
 PR 21-FEB-1991; 91US-0659113.
 PR 21-FEB-1991; 91US-0659114.
 PR 21-FEB-1991; 91US-0659980.
 PR 21-FEB-1991; 91US-0659981.
 PR 14-AUG-1991; 91US-0744870.
 PR 14-AUG-1991; 91US-0745215.
 PR 06-NOV-1991; 91US-0787921.
 XX PA (GILE-) GILEAD SCI INC.
 XX PI Bock LC, Griffin LC, Krawczyk S, Latham JA, Toole JJ;
 PI Muenchau DD;
 XX DR WPI; 1992-316194/38.
 XX PS DNA aptamers specifically binding target molecules - useful for
 PT retrieving target molecules, delivering drugs or toxins to
 PT desired targets and for treating auto-immune diseases
 XX Disclosure; Page 119; 183pp; English.
 XX CC The sequences given in AAQ28473-78 are aptamers which are based on the
 CC unmodified thrombin aptamer given in AAQ28472. These aptamers bind
 CC to thrombin inhibiting its activity, except for the aptamer sequence
 CC given in AAQ28475 which was required in very large quantities to inhibit
 CC thrombin activity. These aptamers are stable, versatile and highly
 CC specific to their intended targets. They can be used to deliver
 CC auxiliary substances, e.g. drugs, toxins, radio isotopes etc. to a
 CC specific part of the body. The aptamers have a binding region of
 CC approx. 10 nucleotide residues.
 XX SQ Sequence 15 BP; 0 A; 0 C; 9 G; 5 T; 0 other;
 Query Match 100.0%; Score 15; DB 13; Length 15;
 Best Local Similarity 93.3%; Pred. No. 4.4e+02;
 Matches 14; Conservative 1; Mismatches 0; Indels 0; Gaps 0;
 QY 1 ggttgggtgggtgg 15
 Db 1 gggttgggtggtggugg 15
 RESULT 5
 AAQ28475
 ID AAQ28475 standard; DNA; 15 BP.
 XX
 AC AAQ28475;
 XX DT 16-FEB-1993 (first entry)
 XX DE Modified thrombin aptamer #3.
 XX KW Aptamer; specifically binding oligonucleotides; primer/linker; PCR;
 KW cleavage; ss.
 XX OS Synthetic.
 XX Key Location/Qualifiers
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 FT misc_difference 2..3
 FT /*tag= b
 FT /note= "The linkage between G2 and T3 is a
 FT misc_difference 3..4
 FT /*tag= c
 FT /note= "The linkage between T3 and T4 is a
 FT misc_difference 4..5
 FT /*tag= d
 FT /note= "The linkage between T4 and G5 is a
 FT misc_difference 5..6
 FT /note= "The linkage between G1 and G2 is a
 FT misc_difference 6..7
 FT /*tag= e
 FT /note= "The linkage between G3 and G4 is a
 FT misc_difference 7..8
 FT /*tag= f
 FT /note= "The linkage between G5 and G6 is a
 FT misc_difference 8..9
 FT /*tag= g
 FT /note= "The linkage between G7 and G8 is a
 FT misc_difference 9..10
 FT /*tag= h
 FT /note= "The linkage between G9 and G10 is a
 FT misc_difference 10..11
 FT /*tag= i
 FT /note= "The linkage between G11 and G12 is a
 FT misc_difference 11..12
 FT /*tag= j
 FT /note= "The linkage between G13 and G14 is a
 FT misc_difference 12..13
 FT /*tag= k
 FT /note= "The linkage between G15 and G16 is a
 FT misc_difference 13..14
 FT /*tag= l
 FT /note= "The linkage between G17 and G18 is a
 FT misc_difference 14..15
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 FT /note= "The linkage between G19 and G20 is a
 FT misc_difference 15..16
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 FT misc_difference 16..17
 FT /*tag= o
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 FT misc_difference 17..18
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 FT /note= "The linkage between G25 and G26 is a
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 FT misc_difference 19..20
 FT /*tag= r
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 FT misc_difference 20..21
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 FT misc_difference 21..22
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 FT /note= "The linkage between G33 and G34 is a
 FT misc_difference 22..23
 FT /*tag= u
 FT /note= "The linkage between G35 and G36 is a
 FT misc_difference 23..24
 FT /*tag= v
 FT /note= "The linkage between G37 and G38 is a
 FT misc_difference 24..25
 FT /*tag= w
 FT /note= "The linkage between G39 and G40 is a
 FT misc_difference 25..26
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 FT misc_difference 26..27
 FT /*tag= y
 FT /note= "The linkage between G43 and G44 is a
 FT misc_difference 27..28
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 FT /note= "The linkage between G45 and G46 is a
 FT misc_difference 28..29
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 FT /note= "The linkage between G47 and G48 is a
 FT misc_difference 29..30
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 FT /note= "The linkage between G49 and G50 is a
 FT misc_difference 30..31
 FT /*tag= cc
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 FT misc_difference 31..32
 FT /*tag= dd
 FT /note= "The linkage between G53 and G54 is a
 FT misc_difference 32..33
 FT /*tag= ee
 FT /note= "The linkage between G55 and G56 is a
 FT misc_difference 33..34
 FT /*tag= ff
 FT /note= "The linkage between G57 and G58 is a
 FT misc_difference 34..35
 FT /*tag= gg
 FT /note= "The linkage between G59 and G60 is a
 FT misc_difference 35..36
 FT /*tag= hh
 FT /note= "The linkage between G61 and G62 is a
 FT misc_difference 36..37
 FT /*tag= ii
 FT /note= "The linkage between G63 and G64 is a
 FT misc_difference 37..38
 FT /*tag= jj
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 FT misc_difference 38..39
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 FT /note= "The linkage between G67 and G68 is a
 FT misc_difference 39..40
 FT /*tag= ll
 FT /note= "The linkage between G69 and G70 is a
 FT misc_difference 40..41
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 FT /note= "The linkage between G71 and G72 is a
 FT misc_difference 41..42
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 FT /note= "The linkage between G73 and G74 is a
 FT misc_difference 42..43
 FT /*tag= oo
 FT /note= "The linkage between G75 and G76 is a
 FT misc_difference 43..44
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 FT /note= "The linkage between G77 and G78 is a
 FT misc_difference 44..45
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 FT /note= "The linkage between G79 and G80 is a
 FT misc_difference 45..46
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 FT /note= "The linkage between G81 and G82 is a
 FT misc_difference 46..47
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 FT misc_difference 47..48
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 FT /note= "The linkage between G85 and G86 is a
 FT misc_difference 48..49
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 FT misc_difference 49..50
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 FT misc_difference 51..52
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 FT misc_difference 52..53
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 FT misc_difference 53..54
 FT /*tag= zz
 FT /note= "The linkage between G97 and G98 is a
 FT misc_difference 54..55
 FT /*tag= aa
 FT /note= "The linkage between G99 and G100 is a
 FT misc_difference 55..56
 FT /*tag= bb
 FT /note= "The linkage between G101 and G102 is a
 FT misc_difference 56..57
 FT /*tag= cc
 FT /note= "The linkage between G103 and G104 is a
 FT misc_difference 57..58
 FT /*tag= dd
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 FT misc_difference 58..59
 FT /*tag= ee
 FT /note= "The linkage between G107 and G108 is a
 FT misc_difference 59..60
 FT /*tag= ff
 FT /note= "The linkage between G109 and G110 is a
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 FT misc_difference 62..63
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 FT misc_difference 63..64
 FT /*tag= jj
 FT /note= "The linkage between G117 and G118 is a
 FT misc_difference 64..65
 FT /*tag= kk
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 FT /note= "The linkage between G121 and G122 is a
 FT misc_difference 66..67
 FT /*tag= mm
 FT /note= "The linkage between G123 and G124 is a
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 FT /*tag= nn
 FT /note= "The linkage between G125 and G126 is a
 FT misc_difference 68..69
 FT /*tag= oo
 FT /note= "The linkage between G127 and G128 is a
 FT misc_difference 69..70
 FT /*tag= pp
 FT /note= "The linkage between G129 and G130 is a
 FT misc_difference 70..71
 FT /*tag= qq
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 FT misc_difference 71..72
 FT /*tag= rr
 FT /note= "The linkage between G133 and G134 is a
 FT misc_difference 72..73
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 FT /note= "The linkage between G141 and G142 is a
 FT misc_difference 76..77
 FT /*tag= ww
 FT /note= "The linkage between G143 and G144 is a
 FT misc_difference 77..78
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 FT /note= "The linkage between G145 and G146 is a
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 FT /note= "The linkage between G147 and G148 is a
 FT misc_difference 79..80
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 FT /note= "The linkage between G149 and G150 is a
 FT misc_difference 80..81
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 FT /note= "The linkage between G151 and G152 is a
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 FT /*tag= bb
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 FT /note= "The linkage between G159 and G160 is a
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 FT /note= "The linkage between G161 and G162 is a
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 FT /note= "The linkage between G163 and G164 is a
 FT misc_difference 87..88
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 FT /note= "The linkage between G165 and G166 is a
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 FT /*tag= vv
 FT /note= "The linkage between G193 and G194 is a
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 FT /*tag= ww
 FT /note= "The linkage between G195 and G196 is a
 FT misc_difference 103..104
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 FT /note= "The linkage between G197 and G198 is a
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 FT /note= "The linkage between G199 and G200 is a
 FT misc_difference 105..106
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 FT /note= "The linkage between G201 and G202 is a
 FT misc_difference 106..107
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 FT /note= "The linkage between G203 and G204 is a
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 FT /*tag= bb
 FT /note= "The linkage between G205 and G206 is a
 FT misc_difference 108..109
 FT /*tag= cc
 FT /note= "The linkage between G207 and G208 is a
 FT misc_difference 109..110
 FT /*tag= dd
 FT /note= "The linkage between G209 and G210 is a
 FT misc_difference 110..111
 FT /*tag= ee
 FT /note= "The linkage between G211 and G212 is a
 FT misc_difference 111..112
 FT /*tag= ff
 FT /note= "The linkage between G213 and G214 is a
 FT misc_difference 112..113
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 FT /note= "The linkage between G215 and G216 is a
 FT misc_difference 113..114
 FT /*tag= hh
 FT /note= "The linkage between G217 and G218 is a
 FT misc_difference 114..115
 FT /*tag= ii
 FT /note= "The linkage between G219 and G220 is a
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 FT /note= "The linkage between G223 and G224 is a
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 FT /note= "The linkage between G225 and G226 is a
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 FT /note= "The linkage between G229 and G230 is a
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 FT /*tag= oo
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 FT /*tag= pp
 FT /note= "The linkage between G233 and G234 is a
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 FT /*tag= qq
 FT /note= "The linkage between G235 and G236 is a
 FT misc_difference 123..124
 FT /*tag= rr
 FT /note= "The linkage between G237 and G238 is a
 FT misc_difference 124..125
 FT /*tag= ss
 FT /note= "The linkage between G239 and G240 is a
 FT misc_difference 125..126
 FT /*tag= tt
 FT /note= "The linkage between G241 and G242 is a
 FT misc_difference 126..127
 FT /*tag= uu
 FT /note= "The linkage between G243 and G244 is a
 FT misc_difference 127..128
 FT /*tag= vv
 FT /note= "The linkage between G245 and G246 is a
 FT misc_difference 128..129
 FT /*tag= ww
 FT /note= "The linkage between G247 and G248 is a
 FT misc_difference 129..130
 FT /*tag= xx
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 FT misc_difference 130..131
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 FT /note= "The linkage between G251 and G252 is a
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 FT /note= "The linkage between G253 and G254 is a
 FT misc_difference 132..133
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 FT /note= "The linkage between G255 and G256 is a
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 FT /note= "The linkage between G257 and G258 is a
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 FT misc_difference 139..140
 FT /*tag= hh
 FT /note= "The linkage between G269 and G270 is a
 FT misc_difference 140..141
 FT /*tag= ii
 FT /note= "The linkage between G271 and G272 is a
 FT misc_difference 141..142
 FT /*tag= jj
 FT /note= "The linkage between G273 and G274 is a
 FT misc_difference 142..143
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 FT /note= "The linkage between G275 and G276 is a
 FT misc_difference 143..144
 FT /*tag= ll
 FT /note= "The linkage between G277 and G278 is a
 FT misc_difference 144..145
 FT /*tag= mm
 FT /note= "The linkage between G279 and G280 is a
 FT misc_difference 145..146
 FT /*tag= nn
 FT /note= "The linkage between G281 and G282 is a
 FT misc_difference 146..147
 FT /*tag= oo
 FT /note= "The linkage between G283 and G284 is a
 FT misc_difference 147..148
 FT /*tag= pp
 FT /note= "The linkage between G285 and G286 is a
 FT misc_difference 148..149
 FT /*tag= qq
 FT /note= "The linkage between G287 and G288 is a
 FT misc_difference 149..150
 FT /*tag= rr
 FT /note= "The linkage between G289 and G290 is a
 FT misc_difference 150..151
 FT /*tag= ss
 FT /note= "The linkage between G291 and G292 is a
 FT misc_difference 151..152
 FT /*tag= tt
 FT /note= "The linkage between G293 and G294 is a
 FT misc_difference 152..153
 FT /*tag= uu
 FT /note= "The linkage between G295 and G296 is a
 FT misc_difference 153..154
 FT /*tag= vv
 FT /note= "The linkage between G297 and G298 is a
 FT misc_difference 154..155
 FT /*tag= ww
 FT /note= "The linkage between G299 and G300 is a
 FT misc_difference 155..156
 FT /*tag= xx
 FT /note= "The linkage between G301 and G302 is a
 FT misc_difference 156..157
 FT /*tag= yy
 FT /note= "The linkage between G303 and G304 is a
 FT misc_difference 157..158
 FT /*tag= zz
 FT /note= "The linkage between G305 and G306 is a
 FT misc_difference 158..159
 FT /*tag= aa
 FT /note= "The linkage between G307 and G308 is a
 FT misc_difference 159..160
 FT /*tag= bb
 FT /note= "The linkage between G309 and G310 is a
 FT misc_difference 160..161
 FT /*tag= cc
 FT /note= "The linkage between G311 and G312 is a
 FT misc_difference 161..162
 FT /*tag= dd
 FT /note= "The linkage between G313 and G314 is a
 FT misc_difference 162..163
 FT /*tag= ee
 FT /note= "The linkage between G315 and G316 is a
 FT misc_difference 163..164
 FT /*tag= ff
 FT /note= "The linkage between G317 and G318 is a
 FT misc_difference 164..165
 FT /*tag= gg
 FT /note= "The linkage between G319 and G320 is a
 FT misc_difference 165..166
 FT /*tag= hh
 FT /note= "The linkage between G321 and G322 is a
 FT misc_difference 166..167
 FT /*tag= ii
 FT /note= "The linkage between G323 and G324 is a
 FT misc_difference 167..168
 FT /*tag= jj
 FT /note= "The linkage between G325 and G326 is a
 FT misc_difference 168..169
 FT /*tag= kk
 FT /note= "The linkage between G327 and G328 is a
 FT misc_difference 169..170
 FT /*tag= ll
 FT /note= "The linkage between G329 and G330 is a
 FT misc_difference 170..171
 FT /*tag= mm
 FT /note= "The linkage between G331 and G332 is a
 FT misc_difference 171..172
 FT /*tag= nn
 FT /note= "The linkage between G333 and G334 is a
 FT misc_difference 172..173
 FT /*tag= oo
 FT /note= "The linkage between G335 and G336 is a
 FT misc_difference 173..174
 FT /*tag= pp
 FT /note= "The linkage between G337 and G338 is a
 FT misc_difference 174..175
 FT /*tag= qq
 FT /note= "The linkage between G339 and G340 is a
 FT misc_difference 175..176
 FT /*tag= rr
 FT /note= "The linkage between G341 and G342 is a
 FT misc_difference 176..177
 FT /*tag= ss
 FT /note= "The linkage between G343 and G344 is a
 FT misc_difference 177..178
 FT /*tag= tt
 FT /note= "The linkage between G345 and G346 is a
 FT misc_difference 178..179
 FT /*tag= uu
 FT /note= "The linkage between G347 and G348 is a
 FT misc_difference 179..180
 FT /*tag= vv
 FT /note= "The linkage between G349 and G350 is a
 FT misc_difference 180..181
 FT /*tag= ww
 FT /note= "The linkage between G351 and G352 is a
 FT misc_difference 181..182
 FT /*tag= xx
 FT /note= "The linkage between G353 and G354 is a
 FT misc_difference 182..183
 FT /*tag= yy
 FT /note= "The linkage between G355 and G356 is a
 FT misc_difference 183..184
 FT /*tag= zz
 FT /note= "The linkage between G357 and G358 is a
 FT misc_difference 184..185
 FT /*tag= aa
 FT /note= "The linkage between G359 and G360 is a
 FT misc_difference 185..186
 FT /*tag= bb
 FT /note= "The linkage between G361 and G362 is a
 FT misc_difference 186..187
 FT /*tag= cc
 FT /note= "The linkage between G363 and G364 is a
 FT misc_difference 187..188
 FT /*tag= dd
 FT /note= "The linkage between G365 and G366 is a
 FT misc_difference 188..189
 FT /*tag= ee
 FT /note= "The linkage between G367 and G368 is a
 FT misc_difference 189..190
 FT /*tag= ff
 FT /note= "The linkage between G369 and G370 is a
 FT misc_difference 190..191
 FT /*tag= gg
 FT /note= "The linkage between G371 and G372 is a
 FT misc_difference 191..192
 FT /*tag= hh
 FT /note= "The linkage between G373 and G374 is a
 FT misc_difference 192..193
 FT /*tag= ii
 FT /note= "The linkage between G375 and G376 is a
 FT misc_difference 193..194
 FT /*tag= jj
 FT /note= "The linkage between G377 and G378 is a
 FT misc_difference 194..195
 FT /*tag= kk
 FT /note= "The linkage between G379 and G380 is a
 FT misc_difference 195..196
 FT /*tag= ll
 FT /note= "The linkage between G381 and G382 is a
 FT misc_difference 196..197
 FT /*tag= mm
 FT /note= "The linkage between G383 and G384 is a
 FT misc_difference 197..198
 FT /*tag= nn
 FT /note= "The linkage between G385 and G386 is a
 FT misc_difference 198..199
 FT /*tag= oo
 FT /note= "The linkage between G387 and G388 is a
 FT misc_difference 199..200

CC unmodified thrombin aptamer given in AAQ28472. These aptamers bind to thrombin inhibiting its activity, except for the aptamer sequence given in AAQ28475 which was required in very large quantities to inhibit thrombin activity. These aptamers are stable, versatile and highly specific to their intended targets. They can be used to deliver auxiliary substances, eg. drugs, toxins, radio isotopes etc. to a specific part of the body. The aptamers have a binding region of approx. 10 nucleotide residues.

CC Sequence 15 BP; 0 A; 0 C; 9 G; 4 T; 0 other;

SQ

Query Match 100.0%; Score 15; DB 13; Length 15;
Best Local Similarity 86.7%; Pred. No. 4.4e+02;
Matches 13; Conservative 2; Mismatches 0; Indels 0; Gaps 0;

QY 1 ggtaggtgtgtttgg 15
|||:||||:|||||

Db 1 ggtaggguguggttgg 15

QULT

7

AAQ28479

standard; DNA; 15 BP.

XX

AC AAQ28479;

XX

DT 16-FEB-1993 (first entry)

XX

DE Modified thrombin aptamer #7.

XX

FH Aptamer; specifically binding oligonucleotides; primer/linker; PCR;

XX

KW cleavage; ss.

XX

OS Synthetic.

XX

PH Location/Qualifiers

FT 3..4

FT misc_difference

/itag= a

/note= "The linkage between T3 and T4 is a formacetal internucleotide linkage"

FT

XX

PN W09214843-A.

XX

PD 03-SEP-1992.

XX

PF 21-FEB-1992;

92WO-US01383.

XX

PR 21-FEB-1991;

91US-06589849.

XX

PR 21-FEB-1991;

91US-0659103.

XX

PR 21-FEB-1991;

91US-0659113.

XX

PR 21-FEB-1991;

91US-0659114.

XX

PR 21-FEB-1991;

91US-0659980.

XX

PR 21-FEB-1991;

91US-0659981.

XX

PR 21-FEB-1991;

91US-0744870.

XX

PR 14-AUG-1991;

91US-0745215.

XX

PR 06-NOV-1991;

91US-0787921.

XX

PA (GILE-) GILEAD SCI INC.

XX

PI Bock LC, Griffin LC, Krawczyk S, Latham JA, Toole JJ;

PI Muenchau DD;

XX

WPI: 1992-316194/38.

XX

DR 1992-316194/38.

XX

PT DNA aptamers specifically binding target molecules - useful for retrieving target molecules, delivering drugs or toxins to

PT desired targets and for treating auto-immune diseases

XX

PS Disclosure; Page 121; 183pp; English.

XX

CC The sequences given in AAQ28479-81 are aptamers which are based on the unmodified thrombin aptamer given in AAQ28472. These aptamers all

CC contain at least one formacetal internucleotide linkages. These CC aptamers bind to thrombin inhibiting its activity. These aptamers CC are stable, versatile and highly specific to their intended targets. CC They can be used to deliver auxiliary substances eg. drugs, toxins, CC radio isotopes etc. to a specific part of the body. The aptamers have CC a binding region of approx. 10 nucleotide residues.

CC Sequence 15 BP; 0 A; 0 C; 9 G; 6 T; 0 other;

SQ

Query Match 100.0%; Score 15; DB 13; Length 15;
Best Local Similarity 100.0%; Pred. No. 4.4e+02;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 ggtaggtgtgtttgg 15
|||:|||||

Db 1 ggtaggguguggttgg 15

RESULT

8

AAQ28480

standard; DNA; 15 BP.

XX

AC AAQ28480;

XX

DT 16-FEB-1993 (first entry)

XX

DE Modified thrombin aptamer #8.

XX

FH Aptamer; specifically binding oligonucleotides; primer/linker; PCR;

XX

KW cleavage; ss.

XX

OS Synthetic.

XX

PH Location/Qualifiers

FT 12..13

FT misc_difference

/itag= a

/note= "The linkage between T12 and T13 is a formacetal internucleotide linkage"

FT

XX

PN W09214843-A.

XX

PD 03-SEP-1992.

XX

PF 21-FEB-1992;

92WO-US01383.

XX

PR 21-FEB-1991;

91US-065896.

XX

PR 21-FEB-1991;

91US-0659849.

XX

PR 21-FEB-1991;

91US-0659103.

XX

PR 21-FEB-1991;

91US-0659113.

XX

PR 21-FEB-1991;

91US-0659114.

XX

PR 21-FEB-1991;

91US-0659980.

XX

PR 21-FEB-1991;

91US-0659981.

XX

PR 14-AUG-1991;

91US-0744870.

XX

PR 14-AUG-1991;

91US-0745215.

XX

PR 06-NOV-1991;

91US-0787921.

XX

PA (GILE-) GILEAD SCI INC.

XX

PI Bock LC, Griffin LC, Krawczyk S, Latham JA, Toole JJ;

PI Muenchau DD;

XX

WPI: 1992-316194/38.

XX

DR 1992-316194/38.

XX

PT DNA aptamers specifically binding target molecules - useful for retrieving target molecules, delivering drugs or toxins to

PT desired targets and for treating auto-immune diseases

XX

PS Disclosure; Page 121; 183pp; English.

XX

CC The sequences given in AAQ28479-81 are aptamers which are based on the unmodified thrombin aptamer given in AAQ28472. These aptamers all

DB 1 ||||||| 15
 1 99ttgggtgtgttgg 15

RESULT 12
 ID AAT28615
 XX AAT28615 standard; DNA; 15 BP.

DE AAT28615
 XX
 DT 08-MAY-1998 (first entry)
 XX Spectroscopically detectable nucleic acid ligand compound #1.
 KW
 fluorescein; thiazole orange; ss.

OS Synthetic.

XX

KEY modified_base Location/Qualifiers

FT 1 /**tag= a
 /note= "Guanine6 and Thymine7 optionally linked via
 a C6 linker molecule or labeled with Thiazole
 Orange via a C3 linker"

FT misc_feature 6..7 /**tag= b
 /note= "Thymine6 and Thymine7 optionally linked via
 a phosphorothioate bond"
 linked to fluorescein via iodoacetamide."

FT modified_base 7 /**tag= c
 /note= "Thymine6 phosphorothioate when present is
 linked to fluorescein via iodoacetamide."

XX WO9622383-A1.

PD 25-JUL-1996.
 XX
 PR 21-JUL-1995; 95WO-US09237.

XX 20-JAN-1995; 95US-0376329.
 PR (BECT) BECTON DICKINSON CO.
 PA (NEXS-) NEXSTAR PHARM INC.

XX Gold L, Malinowski DP, Pitner JB, Vonk GP;
 PT XX DR WPI; 1996-35454/35.

XX
 PT Detection of target cpds. such as thrombin - using spectroscopically
 detectable nucleic acid ligands

XX
 PS Claims 7 and 11; Figure 1; 37pp; English.

XX
 CC This sequence represents spectroscopically detectable nucleic acid
 ligands which were used to detect the presence or absence of a target
 compound (thrombin) in a sample. This ligand can also be used for
 monitoring the binding of target compounds (such as growth factors) to
 their receptors in competition-based assays. It is thus useful in
 diagnostic assays. Spectroscopically detectable nucleic acid
 ligands of the invention may be used to detect e.g. thrombin, elastase,
 cell surface markers, growth factors, receptor receptors, whole
 cells or viral particles present in biological samples such as blood.
 The receptor molecules are typically relatively small in relation to
 traditional receptor molecules such as antibodies. Any additional weight
 or volume added to the receptor molecules (even in the form of a small
 target) will significantly increase the weight or volume of the
 labelled receptor molecule and therefore permit detection of the
 relatively significant energy differences involved (between bound and
 unbound labelled receptor molecules).

XX SQ Sequence 15 BP; 0 A; 0 C; 9 G; 6 T; 0 other;

Query Match 100.0%; Score 15; DB 17; Length 15;
 Best Local Similarity 100.0%; Pred. No. 4.4e+02;
 Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0; Qy 1 ggttgggtgtgttgg 15

DB 10..11
 1 99ttgggtgtgttgg 15

RESULT 12
 ID AAT28615
 XX AAT28615 standard; DNA; 15 BP.

DE AAT28615
 XX
 DT 14-NOV-1996 (first entry)
 XX Quadruplex/duplex thrombin inhibitor consensus sequence #3.
 KW thrombin; inhibitor; quadruplex/duplex structure; multimeric compound;
 KW quadruplex motif; telomerase; nuclease resistant; diagnostic probe;
 KW therapy; ss.

OS Synthetic.

KEY misc_feature Location/Qualifiers

FT 1..2 /**tag= a
 /note= "bases involved in quadruplex formation"

FT misc_feature 4 /**tag= b
 /note= "forms base pair with T at position 13"

FT misc_feature 5..6 /**tag= c
 /note= "bases involved in quadruplex formation"

FT misc_feature 10..11 /**tag= d
 /note= "bases involved in quadruplex formation"

FT misc_feature 13 /**tag= e
 /note= "forms base pair with T at position 4"

FT misc_feature 14..15 /**tag= f
 /note= "bases involved in quadruplex formation"

XX WO9611010-A1.

PD 18-APR-1996.
 XX
 PR 20-SEP-1995; 95WO-US11985.

XX 07-OCT-1994; 94US-0320139.
 PR
 XX
 PA (PHAR) PHARMAGENICS INC.

XX
 PT Bertelsen AH, Beutel BA, Cook AF, Gao H, Joesten ME;
 PT Macaya RF;
 XX DR WPI; 1996-209651/21.

XX
 PT Single stranded oligo:deoxy:ribonucleotide thrombin inhibitors -
 PT comprise quadruplex consensus motif either flanked by complementary
 PT sequences which form duplex-stem or having bridged termini

XX PS Example; Page 41; 61pp; English.

XX
 CC AAT28613-n28615, and AAT28625 represent consensus sequences for the
 CC thrombin inhibitors of the invention. These sequences can be used to
 CC inhibit and to detect thrombin activity. These sequences form
 CC quadruplex/duplex structures, and can also form multimeric compounds.
 CC The inhibitors of the invention bind thrombin with a higher affinity than
 CC quadruplex structures that lack the duplex stem (such as this sequence).
 CC By bridging the 5' and 3' ends of these structures may optimise their use
 CC as inhibitors to targets other than thrombin, such as telomerase. The
 CC secondary structures formed by these sequences result in molecules which
 CC are constrained to the most biologically active conformation. The
 CC bridged molecules are up to 45 times more nuclease resistant than the
 CC unbridged molecules. These sequences can be used as diagnostic probes

CC to monitor the presence of thrombin, and thereby determine whether there
 CC is a need to modulate its function or activity. The inhibitors can also
 CC be administered to a cell in order to prevent the deleterious
 CC consequences of overproduction of thrombin, or to effect the benefits of
 CC inhibition of thrombin function.

SQ

Sequence 15 BP; 0 A; 0 C; 9 G; 6 T; 0 other;

SQ

Query Match 100.0%; Score 15; DB 17; Length 15;

Best Local Similarity 100.0%; Pred. No. 4.4e+02; Mismatches 0; Indels 0; Gaps 0;
 Matches 15; Conservative 0; Misnatches 0; Indels 0; Gaps 0;

Qy 1 ggttgggtgggtgg 15
 Db 1 ggttgggtgggtgg 15

SQ

Sequence 15 BP; 0 A; 0 C; 9 G; 6 T; 0 other;

SQ

RESULT 13

1/7808

AAT17808 standard; DNA; 15 BP.

XX

AAT17808;

XX

DT 30-OCT-1996 (first entry)

XX

DE Glycosaminoglycan-degrading enzyme inhibitor IgLPS.

XX

KW heparanase; heparitinase; mammalian; bacterial; platelet; macrophage;

KW neutrophil; leukocyte; endothelial cell; smooth muscle cell; carcinoma;

KW tumour cell; activation; proliferation; migration; cancer; inflammation;

KW autoimmune disorder; infection; pathogenic organism; atherosclerosis;

KW cardiovascular disease; vascular hyperplasia; restenosis; therapy; ss.

OS Synthetic.

XX

FH key modified_base Location/Qualifiers

FT 1..15

FT /*tag= a

FT /note= "optionally phosphorothioated, or phosphorodithioated backbone"

XX

W09608559-A1.

XX

PD 21-MAR-1996.

XX

PA 13-SEP-1995; 95WO-AU00600.

XX

PR 14-AUG-1995; 95AU-0004769.

PR 16-SEP-1994; 94AU-0008226.

PR 16-SEP-1994; 94AU-0008227.

PA (CARD-) CARDIAC CRC NOMINEES PTY LTD.

XX

PT Graham L, Underwood PA;

XX

DR WPI; 1996-179936/18.

XX

PT Oligo:nucleotide(s) having sulphur substituents between nucleoside(s) - for inhibiting glycosaminoglycan-degrading enzymes, for treating, e.g. cancer, inflammation, infection or autoimmune disorders.

PT

Example 2: Page 33: 73pp: English.

XX

AAT17805-117808, and AAT17810-17813 represent glycosaminoglycan-degrading enzyme (GDE) inhibitors. The GDEs which

CC

these sequences inhibit are endo-lycosidases (which cleave known as heparinases) of mammalian or bacterial origin. These

CC

sequences can be used for inhibiting GDEs associated with platelets, macrophages, neutrophils, leukocytes, endothelial cells, smooth muscle

CC

cells, carcinoma and tumour cells, and bacteria. They can also be used

CC to inhibit smooth muscle cell activation, proliferation or migration.
 CC The sequences can be used to treat cancer, inflammation, autoimmune
 CC disorders, infection caused by pathogenic organisms, and cardiovascular
 CC disease, such as vascular hyperplasia, restenosis and atherosclerosis.
 CC These inhibitors can also be used as biochemical reagents for studying
 CC GDE activities and mechanisms of enzyme activity.

SQ

Query Match 100.0%; Score 15; DB 17; Length 15;

Best Local Similarity 100.0%; Pred. No. 4.4e+02; Mismatches 0; Indels 0; Gaps 0;
 Matches 15; Conservative 0; Misnatches 0; Indels 0; Gaps 0;

Qy 1 ggttgggtgtgtgg 15
 Db 1 ggttgggtgtgtgg 15

SQ

Sequence 15 BP; 0 A; 0 C; 9 G; 6 T; 0 other;

SQ

RESULT 14

AAT51669

ID AAT51669 standard; DNA; 15 BP.

XX

AAT51669;

XX

DT 12-NOV-1997 (first entry)

XX

DE Viral integrase inhibiting oligonucleotide.

XX

KW Human immunodeficiency virus; HIV; Epstein Barr virus; EBV; herpes simplex virus; HSV; human papilloma virus; HPV; adebovirus; respiratory syncytial virus; RSV; cytomegalovirus; CMV; hepatitis B; integrase inhibition; guanosine tetrad; ss.

XX

OS Synthetic.

XX

FH key modified_base Location/Qualifiers

FT 1..15

FT /*tag= a

FT /note= "optionally contains all phosphorothioate linkages or a phosphorothioate linkage between penultimate and last nucleotide at 3' end"

FT

FT

W09703997-A1.

XX

PD 06-FEB-1997.

XX

PR 17-JUL-1996; 96WO-US11786.

XX

PR 23-APR-1996; 96US-0016271.

XX

PR 19-JUL-1995; 95US-0001505.

XX

PR 23-OCT-1995; 95US-0535160.

XX

PR 19-MAR-1996; 96US-0013688.

XX

PR 25-MAR-1996; 96US-0014007.

XX

PR 17-APR-1996; 96US-0015714.

XX

PA (ARON-) ARONEX PHARM INC.

XX

PT Fennewald S, Hogan ME, Mazumder A, Ojwang JO, Pommier Y;

PT

PT Rando RF, Zendejas JG;

XX

DR WPI; 1997-132569/12.

XX

PT Oligo:nucleotide(s) capable of forming guanosine tetrads - inhibit viral enzyme responsible for integrating viral nucleic acid into the

PT

PT host genome

XX

PS Claim 3; Page 81; 245pp: English.

XX

CC AAT51619-T51628 are oligonucleotides used to inhibit the production of viruses within a host cell. The oligonucleotides may form guanosine tetrads (structures formed of eight hydrogen bonds by coordination of

CC

CC to the centre of a quadruplex, and by strong stacking interactions) and CC are used to prevent the integration of viral nucleic acid into a host CC genome. The oligonucleotides inhibit functioning of the integrase enzyme CC and hence prevent viral infection. Viral infections that may be treated CC include human immunodeficiency virus (HIV), Epstein Barr virus (EBV), CC herpes simplex virus (HSV), human papilloma virus (HPV), adenovirus, CC respiratory syncytial virus (RSV), cytomegalovirus (CMV) and hepatitis B virus (HBV), especially HIV-1 infection.

SQ Sequence 15 BP; 0 A; 0 C; 9 G; 6 T; 0 other;

Query Match 100.0%; Score 15; DB 18; Length 15;

Best Local Similarity 100.0%; Pred. No. 4.4e+02; Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 gggtgggtgggtgg 15

Db 1 gggtgggtgggtgg 15

SQI 15
RAT8811
ID AAT85811 standard; DNA; 15 BP.

XX
XX
AC AAT85811;
XX
XX
DT 11-NOV-1997 (first entry)
XX
DE Thrombin-binding nucleic acid ligand.
XX
KW Thiazole orange; fluorescein; spectroscopic assay;
XX fluorescence polarisation detection; thrombin; ss.
OS Synthetic.

XX
FH Location/Qualifiers
FT
FT 1
FT /*tag= a
FT /note= "5'-labelled either with thiazole orange via
FT aminopropyl C3 linker arm (= Compound 3) or
FT with fluorescein via a C6 linker arm
FT (= Compound 1)",
XX
PN US5650275-A.
XX
PD 22-JUL-1997.
XX
XX
PR 18-MAY-1995; 95US-0443957.
PR 11-JUN-1990; 90US-0536428.
PR 10-JUN-1991; 91US-0714131.
PR 17-AUG-1992; 92US-0931473.
PR 07-OCT-1993; 93US-0134028.
PR 28-APR-1994; 94US-0234997.
PR 18-JUL-1994; 94US-0276271.
XX
PA (GOLD/ GOLD L.
PA (MALI/ MALINOWSKI D P.
PA (PITN/ PITNER J B.
PA (VONK/ VONK G P.
PI Gold L, Malinowski DP, Pitner JB, Vonk GP;
XX
DR WPI; 1997-384664/35.

CC Spectroscopically detectable labelled nucleic acid ligands are used CC in a claimed method for determining the presence of a target compound CC in a sample. An increase in the spectroscopic emission of the CC ligand in the presence of a sample relative to the ligand alone is CC indicative of the presence of the target compound in the sample. CC Target molecules may be proteins, peptides, cell surface markers, CC carbohydrates, polysaccharides, glycoproteins, hormones, receptors, CC antigens, antibodies, co-factors, inhibitors, drugs, dyes, nutrients, CC growth factors, amino acids, ATP, whole cells or viral particles. CC The present sequence is a preferred nucleic acid ligand for CC detecting thrombin. When labelled with fluorescein it is designated CC Compound 1 (claim 6) and when labelled with thiazole orange it is CC designated compound 3 (claim 15).

SQ Sequence 15 BP; 0 A; 0 C; 9 G; 6 T; 0 other;

Query Match 100.0%; Score 15; DB 18; Length 15;

Best Local Similarity 100.0%; Pred. No. 4.4e+02; Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 gggtgggtgggtgg 15

Db 1 gggtgggtgggtgg 15

Search completed: June 6, 2002, 16:08:47
Job time: 2148 sec

RESULT 2
 US-08-219-012-1
 Sequence 1, Application US/08219012
 Patent No. 5,943,293
 GENERAL INFORMATION:
 APPLICANT: Larry Gold
 APPLICANT: Diane Tasset
 TITLE OF INVENTION: Ligands of Thrombin
 NUMBER OF SEQUENCES: 92
 CORRESPONDENCE ADDRESS:
 ADDRESSE: Beaton & Swanson, P.C.
 STREET: 4582 South Ulster Street, Parkway, Suite #403
 CITY: Denver
 STATE: Colorado
 COUNTRY: USA
 ZIP: 80237
 COMPUTER READABLE FORM:
 MEDIUM TYPE: Floppy disk
 COMPUTER: IBM PC compatible
 OPERATING SYSTEM: PC-DOS/MS-DOS
 SOFTWARE: PatentIn Release #1.0, Version #1.25
 CURRENT APPLICATION DATA:
 APPLICATION NUMBER: US/08/376,329
 FILING DATE: 08/08/2001
 CLASSIFICATION: 435
 ATTORNEY/AGENT INFORMATION:
 NAME: Highet, David W
 REGISTRATION NUMBER: 30,265
 REFERENCE/DOCKET NUMBER: P-3126
 TELECOMMUNICATION INFORMATION:
 TELEPHONE: 201 848 9228
 TELEFAX: 201 847 5317
 INFORMATION FOR SEQ ID NO: 1:
 SEQUENCE CHARACTERISTICS:
 LENGTH: 15 base pairs
 TYPE: nucleic acid
 STRANDEDNESS: single
 TOPOLOGY: linear
 CURRENT APPLICATION DATA:
 APPLICATION NUMBER: US/08/219,012
 FILING DATE:
 CLASSIFICATION: 435
 PRIOR APPLICATION DATA: none
 ATTORNEY/AGENT INFORMATION:
 NAME: Barry J. Swanson
 REGISTRATION NUMBER: 33,215
 REFERENCE/DOCKET NUMBER:
 TELECOMMUNICATION INFORMATION:
 TELEPHONE: (303) 850-9900
 TELEFAX: (303) 850-9401
 INFORMATION FOR SEQ ID NO: 1:
 SEQUENCE CHARACTERISTICS:
 LENGTH: 15 base pairs
 TYPE: nucleic acid
 STRANDEDNESS: single
 TOPOLOGY: linear
 RESULT 3
 US-08-376-329-1
 Query Match 100.0%; Score 15; DB 1; Length 15;
 Best Local Similarity 100.0%; Pred. No. 48;
 matches 15; Conservative 0; Mismatches 0;
 QY 1 gggtgggtgggtgg 15
 QY 1 ||||||| 15
 Db 1 GGTGGGGGGGG 15
 RESULT 4
 US-08-276-271-1
 Sequence 1, Application US/08276271
 Patent No. 5,650,275
 GENERAL INFORMATION:
 APPLICANT: Pitner, James B
 APPLICANT: Malinowski, Douglas P
 APPLICANT: Vonk, Glenn P
 APPLICANT: Gold, Larry
 TITLE OF INVENTION: Spectroscopically Detectable Nucleic
 TITLE OF INVENTION: Acid Ligands
 NUMBER OF SEQUENCES: 4
 CORRESPONDENCE ADDRESS:
 ADDRESSE: Richard J. Roderick, Becton Dickinson and
 COMPANY: Company
 STREET: 1 Becton Drive
 CITY: Franklin Lakes
 STATE: NJ
 COUNTRY: USA
 ZIP: 07417-1880
 COMPUTER READABLE FORM:
 MEDIUM TYPE: Floppy disk
 COMPUTER: IBM PC compatible
 OPERATING SYSTEM: PC-DOS/MS-DOS
 SOFTWARE: PatentIn Release #1.0, Version #1.25
 CURRENT APPLICATION DATA:
 APPLICATION NUMBER: US/08/276,271
 FILING DATE:
 CLASSIFICATION: 436
 ATTORNEY/AGENT INFORMATION:
 NAME: Highet, David W
 REGISTRATION NUMBER: 30,265
 REFERENCE/DOCKET NUMBER: P-3126
 TELECOMMUNICATION INFORMATION:
 TELEPHONE: 201 848 9228
 TELEFAX: 201 847 5317
 INFORMATION FOR SEQ ID NO: 1:

SEQUENCE CHARACTERISTICS:
 LENGTH: 15 base pairs
 TYPE: nucleic acid
 STRANDEDNESS: single
 TOPOLOGY: linear
 MOLECULE TYPE: DNA (genomic)

US-08-276-271-1

Query Match Similarity 100.0%; Score 15; DB 1; Length 15;
 Best Local Similarity 100.0%; Pred. No. 48; 0; Mismatches 0;
 Matches 15; Conservative 0; Indels 0; Gaps 0;
 QY 1 gggtgggtgggtgg 15
 DB 1 GGTGGTGGTGG 15

RESULT 5

US-08-539-516-4

Sequence 4, Application US/08539516

PATENT NO. 5668738

GENERAL INFORMATION:

APPLICANT: Nadeau, James G.
 APPLICANT: Ciolekowski, Mary Lee
 APPLICANT: Vogler, Erwin A.
 TITLE OF INVENTION: BI-DIRECTIONAL OLIGONUCLEOTIDES THAT
 BIND PROTEIN
 NUMBER OF SEQUENCES: 4
 CORRESPONDENCE ADDRESS:
 ADDRESSEE: Richard J. Rodrick, Becton Dickinson and
 COMPANY: Company
 STREET: 1 Becton Drive
 CITY: Franklin Lakes
 STATE: NJ
 COUNTRY: US
 ZIP: 07417

COMPUTER READABLE FORM:

COMPUTER TYPE: Floppy disk
 COMPUTER: IBM PC compatible
 OPERATING SYSTEM: PC-DOS/MS-DOS
 SOFTWARE: Patentin Release #1.0, Version #1.25
 CURRENT APPLICATION DATA:
 APPLICATION NUMBER: US/08/614,447

CURRENT APPLICATION DATA:
 APPLICATION NUMBER: US/08/539,516

FILING DATE: 11-DEC-1995

CLASSIFICATION: 435

PRIOR APPLICATION DATA:

APPLICATION NUMBER: US/08/614,447

FILING DATE: 12-MAR-1996

APPLICATION NUMBER: US/08/252,071

FILING DATE: 31-MAY-1994

ATTORNEY/AGENT INFORMATION:

NAME: Hight, David W.
 REGISTRATION NUMBER: 30,265

REFERENCE/DOCKET NUMBER: P-3028

TELECOMMUNICATION INFORMATION:

TELEPHONE: (201) 847-3317

TELEFAX: (201) 848-9228

INFORMATION FOR SEQ ID NO: 4:

SEQUENCE CHARACTERISTICS:

LENGTH: 15 base pairs

TYPE: nucleic acid

STRANDEDNESS: single

TOPOLOGY: linear

MOLECULE TYPE: DNA (genomic)

US-08-539-516-4

Db 1 ||||||| 15

RESULT 6
 Sequence 4, Application US/08614447
 Patent No. 5668738

US-08-614-447-4

GENERAL INFORMATION:
 APPLICANT: Nadeau, James G.
 APPLICANT: Ciolekowski, Mary Lee
 TITLE OF INVENTION: BI-DIRECTIONAL OLIGONUCLEOTIDES THAT
 BIND PROTEIN
 NUMBER OF SEQUENCES: 4
 CORRESPONDENCE ADDRESS:
 ADDRESSEE: Richard J. Rodrick, Becton Dickinson and
 COMPANY: Company
 STREET: 1 Becton Drive
 CITY: Franklin Lakes
 STATE: NJ
 COUNTRY: US
 ZIP: 07417

COMPUTER READABLE FORM:
 COMPUTER TYPE: Floppy disk
 COMPUTER: IBM PC compatible
 OPERATING SYSTEM: PC-DOS/MS-DOS
 SOFTWARE: Patentin Release #1.0, Version #1.25
 CURRENT APPLICATION DATA:
 APPLICATION NUMBER: US/08/614,447
 FILING DATE: 12-MAR-1996
 CLASSIFICATION: 435
 PRIOR APPLICATION DATA:
 APPLICATION NUMBER: US 08/252,071
 APPLICATION NUMBER: US/08/614,447
 FILING DATE: 31-MAY-1994
 ATTORNEY/AGENT INFORMATION:
 NAME: Hight, David W.
 REGISTRATION NUMBER: 30,265
 REFERENCE/DOCKET NUMBER: P-3028
 TELECOMMUNICATION INFORMATION:
 TELEPHONE: (201) 847-3317
 TELEFAX: (201) 848-9228
 INFORMATION FOR SEQ ID NO: 4:
 SEQUENCE CHARACTERISTICS:
 LENGTH: 15 base pairs
 TYPE: nucleic acid
 STRANDEDNESS: single
 TOPOLOGY: linear
 MOLECULE TYPE: DNA (genomic)

RESULT 7

US-08-0703-755A-3

Sequence 3, Application US/08703755A

PATENT NO. 5691145

GENERAL INFORMATION:

APPLICANT: Pittner, Bruce

APPLICANT: Vonk, Glenn P.

APPLICANT: Nadeau, James G.

TITLE OF INVENTION: DIRECTION OF NUCLEIC ACIDS USING

TITLE OF INVENTION: G-QUARTETS

NUMBER OF SEQUENCES: 7

CORRESPONDENCE ADDRESS:

Query Match Similarity 100.0%; Score 15; DB 1; Length 15;
 Best Local Similarity 100.0%; Pred. No. 48; 0; Mismatches 0;
 Matches 15; Conservative 0; Indels 0; Gaps 0;
 QY 1 gggtgggtgggtgg 15
 DB 1 GGTGGTGGTGG 15

QY 1 gggtgggtgggtgg 15

ADDRESSEE: Richard J. Rodrick, Becton Dickinson and Company
 STREET: 1 Becton Drive
 CITY: Franklin Lakes
 STATE: NJ
 COUNTRY: US
 ZIP: 07417
 COMPUTER READABLE FORM:
 COMPUTER: IBM PC compatible
 OPERATING SYSTEM: PC-DOS/MS-DOS
 SOFTWARE: PatentIn Release #1.0, Version #1.30
 CURRENT APPLICATION DATA:
 APPLICATION NUMBER: US/08/703,755A
 FILING DATE:
 CLASSIFICATION: 435
 ATTORNEY/AGENT INFORMATION:
 NAME: Fugit, Donna R.
 REFERENCE/DOCKET NUMBER: 32,135
 INFORMATION FOR SEQ ID NO: 3:
 SEQUENCE CHARACTERISTICS:
 LENGTH: 15 base pairs
 TYPE: nucleic acid
 STRANDEDNESS: single
 TOPOLOGY: linear
 US-08-703-755A-3

RESULT 8
 US-08-484-192-29
 ; Sequence 29, Application US/08484192
 ; Patent No. 5756291
 ; GENERAL INFORMATION:
 ; APPLICANT: GRIFFIN, LINDA C.
 ; APPLICANT: ALBRECHT, GLENN
 ; APPLICANT: LATHAM, JOHN
 ; APPLICANT: LEONG, LAWRENCE
 ; APPLICANT: VERMAAS, ERIC
 ; TITLE OF INVENTION: AMMERS SPECIFIC FOR BIOMOLECULES AND METHODS OF MAKING
 ; NUMBER OF SEQUENCES: 181
 ; CORRESPONDENCE ADDRESS:
 ; ADDRESSEE: MORRISON & FOERSTER
 ; STREET: 755 PAGE MILL ROAD
 ; CITY: PALO ALTO
 ; STATE: CALIFORNIA
 ; COUNTRY: USA
 ; ZIP: 94304
 COMPUTER READABLE FORM:
 COMPUTER: IBM PC compatible
 OPERATING SYSTEM: PC-DOS/MS-DOS
 SOFTWARE: PatentIn Release #1.0, Version #1.25
 CURRENT APPLICATION DATA:
 APPLICATION NUMBER: US/08/484,192
 FILING DATE:
 CLASSIFICATION: 435
 PRIORITY APPLICATION DATA:
 COMPUTER: IBM PC compatible
 OPERATING SYSTEM: PC-DOS/MS-DOS
 SOFTWARE: PatentIn Release #1.0, Version #1.25
 CURRENT APPLICATION DATA:
 APPLICATION NUMBER: US/08/484,192
 FILING DATE:
 CLASSIFICATION: 435
 PRIORITY APPLICATION DATA:
 APPLICATION NUMBER: US 07/934,387
 FILING DATE: 21-AUG-1992
 ATTORNEY/AGENT INFORMATION:
 NAME: GRACEY, NANCY J.
 REFERENCE/DOCKET NUMBER: 246102002221
 TELECOMMUNICATION INFORMATION:
 TELEPHONE: 415-813-5600
 TELEFAX: 415-494-0792
 TELEX: 705141
 INFORMATION FOR SEQ ID NO: 31:
 SEQUENCE CHARACTERISTICS:
 LENGTH: 15 base pairs
 TYPE: nucleic acid
 STRANDEDNESS: single
 TOPOLOGY: linear
 NAME/KEY: misc_difference

RESULT 9
 US-08-484-192-31
 ; Sequence 31, Application US/08484192
 ; Patent No. 5756291
 ; GENERAL INFORMATION:
 ; APPLICANT: GRIFFIN, LINDA C.
 ; APPLICANT: ALBRECHT, GLENN
 ; APPLICANT: LATHAM, JOHN
 ; APPLICANT: LEONG, LAWRENCE
 ; APPLICANT: VERMAAS, ERIC
 ; TITLE OF INVENTION: AMMERS SPECIFIC FOR BIOMOLECULES AND METHODS OF MAKING
 ; NUMBER OF SEQUENCES: 181
 ; CORRESPONDENCE ADDRESS:
 ; ADDRESSEE: MORRISON & FOERSTER
 ; STREET: 755 PAGE MILL ROAD
 ; CITY: PALO ALTO
 ; STATE: CALIFORNIA
 ; COUNTRY: USA
 ; ZIP: 94304
 COMPUTER READABLE FORM:
 COMPUTER: IBM PC compatible
 OPERATING SYSTEM: PC-DOS/MS-DOS
 SOFTWARE: PatentIn Release #1.0, Version #1.25
 CURRENT APPLICATION DATA:
 APPLICATION NUMBER: US/08/484,192
 FILING DATE:
 CLASSIFICATION: 435
 PRIORITY APPLICATION DATA:
 APPLICATION NUMBER: US 07/934,387
 FILING DATE: 21-AUG-1992
 ATTORNEY/AGENT INFORMATION:
 NAME: GRACEY, NANCY J.
 REFERENCE/DOCKET NUMBER: 246102002221
 TELECOMMUNICATION INFORMATION:
 TELEPHONE: 415-813-5600
 TELEFAX: 415-494-0792
 TELEX: 705141
 INFORMATION FOR SEQ ID NO: 31:
 SEQUENCE CHARACTERISTICS:
 LENGTH: 15 base pairs
 TYPE: nucleic acid
 STRANDEDNESS: single
 TOPOLOGY: linear
 NAME/KEY: misc_difference

LOCATION: replace(13..15, "")
 OTHER INFORMATION: /note= "These positions are thioate
 OTHER INFORMATION: (i.e., P(OJS) linked."
 US-08-484-192-31

RESULT 10
 Best Local Similarity 100.0%; Score 15; DB 1; Length 15;
 Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 1 gggtgggtgggtgg 15
 Db 1 GGTTGGTGGTGG 15

RESULT 11
 US-08-484-192-36
 Sequence 36 Application US/08484192
 Patient No. 5756291
 GENERAL INFORMATION:
 APPLICANT: GRIFFIN, LINDA C.
 APPLICANT: ALBRECHT, GLENN
 APPLICANT: LATHAM, JOHN
 APPLICANT: LEUNG, LAWRENCE
 APPLICANT: VERMAAS, ERIC
 APPLICANT: TOOLE, JOHN J.
 TITLE OF INVENTION: APTAMERS SPECIFIC FOR BIOMOLECULES AND
 TITLE OF INVENTION: METHODS OF MAKING
 NUMBER OF SEQUENCES: 181
 CORRESPONDENCE ADDRESS:
 ADDRESSEE: MORRISON & FOERSTER
 STREET: 755 PAGE MILL ROAD
 CITY: PALO ALTO
 STATE: CALIFORNIA
 COUNTRY: USA
 ZIP: 94304
 COMPUTER READABLE FORM:
 COMPUTER TYPE: FLOPPY DISK
 COMPUTER: IBM PC COMPATIBLE
 OPERATING SYSTEM: PC-DOS/MS-DOS
 SOFTWARE: PATENTIN RELEASE #1.0, Version #1.25
 CURRENT APPLICATION DATA:
 APPLICATION NUMBER: US/08/484,192
 FILING DATE:
 CLASSIFICATION: 435
 PRIORITY APPLICATION DATA:
 FILING DATE: 21-AUG-1992
 ATTORNEY/AGENT INFORMATION:
 NAME: GRACEY, NANCY J.
 REGISTRATION NUMBER: 28,216
 REFERENCE/DOCKET NUMBER: 246102002221
 TELECOMMUNICATION INFORMATION:
 TELEPHONE: 415-813-5600
 TELEFAX: 415-494-0792
 TELEX: 706141
 INFORMATION FOR SEQ ID NO: 36:
 SEQUENCE CHARACTERISTICS:
 LENGTH: 15 base pairs
 TYPE: nucleic acid
 STRANDEDNESS: single
 TOPOLOGY: linear
 FEATURE:
 NAME/KEY: misc_difference
 LOCATION: replace(3..4, "")
 OTHER INFORMATION: /note= "This is a formacetal
 OTHER INFORMATION: linkage."
 US-08-484-192-36

Query Match 100.0%; Score 15; DB 1; Length 15;
 Best Local Similarity 100.0%; Score 15; DB 1; Length 15;
 Matches 15; Conservative 0; Mismatches 0; Indels 0;
 QY 1 gggtgggtgggtgg 15
 Db 1 GGTTGGTGGTGG 15

Patent No. 5756291

GENERAL INFORMATION:

APPLICANT: GRIFFIN, LINDA C.

APPLICANT: ALBRECHT, GLENN

APPLICANT: LATHAM, JOHN

APPLICANT: LEUNG, LAWRENCE

APPLICANT: VERMAS, ERIC

APPLICANT: TOOLE, JOHN J.

TITLE OF INVENTION: APTAMERS SPECIFIC FOR BIOMOLECULES AND

NUMBER OF SEQUENCES: 181

CORRESPONDENCE ADDRESS: MORRISON & FOERSTER

STREET: 755 PAGE MILL ROAD

CITY: PALO ALTO

STATE: CALIFORNIA

COUNTRY: USA

ZIP: 94304

COMPUTER READABLE FORM:

COMPUTER: IBM PC compatible

OPERATING SYSTEM: PC-DOS/MS-DOS

SOFTWARE: PatentIn Release #1.0, Version #1.25

CURRENT APPLICATION DATA:

APPLICATION NUMBER: US/08/484,192

FILING DATE: 21-AUG-1992

CLASSIFICATION: 435

PRIORITY APPLICATION DATA:

APPLICATION NUMBER: US/07/934,387

NAME: GRACEY, NANCY J.

REGISTRATION NUMBER: 28,216

REFERENCE/DOCKET NUMBER: 246102002221

TELECOMMUNICATION INFORMATION:

TELEPHONE: 415-813-5600

TELEFAX: 415-494-0792

TELEPAC: 706141

INFORMATION FOR SEQ ID NO: 37:

SEQUENCE CHARACTERISTICS:

LENGTH: 15 base pairs

TYPE: nucleic acid

STRANDEDNESS: single

TOPOLOGY: linear

FEATURE:

NAME/KEY: misc_difference

LOCATION: replace(3..4, "")

OTHER INFORMATION: /note= "This is a formacetal

FEATURE:

NAME/KEY: misc_difference

LOCATION: replace(12..13, "")

OTHER INFORMATION: /note= "This is a formacetal

OTHER INFORMATION: linkage.

US-08-484-192-37

Query Match 100 %; Score 15; DB 1; length 15;

Best Local Similarity 100.0%; Pred. No. 48;

Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

RESULT 13

US-08-484-192-38

Sequence 38, Application US/08484192

; Patent No. 5756291

; GENERAL INFORMATION:

; APPLICANT: GRIFFIN, LINDA C.

; APPLICANT: ALBRECHT, GLENN

; APPLICANT: LATHAM, JOHN

; APPLICANT: LEUNG, LAWRENCE

; APPLICANT: VERMAS, ERIC

; APPLICANT: TOOLE, JOHN J.

; APPLICANT: LATHAM, JOHN

; APPLICANT: LEUNG, LAWRENCE

; APPLICANT: VERMAS, ERIC

; APPLICANT: TOOLE, JOHN J.

TITLE OF INVENTION: APTAMERS SPECIFIC FOR BIOMOLECULES AND

NUMBER OF SEQUENCES: 181

CORRESPONDENCE ADDRESS: MORRISON & FOERSTER

ADRESSE: MORRISON & FOERSTER

STREET: 755 PAGE MILL ROAD

CITY: PALO ALTO

STATE: CALIFORNIA

COUNTRY: USA

ZIP: 94304

COMPUTER READABLE FORM:

COMPUTER: IBM PC compatible

OPERATING SYSTEM: PC-DOS/MS-DOS

SOFTWARE: PatentIn Release #1.0, Version #1.25

CURRENT APPLICATION DATA:

APPLICATION NUMBER: US/08/484,192

FILING DATE: 21-AUG-1992

CLASSIFICATION: 435

PRIORITY APPLICATION DATA:

APPLICATION NUMBER: US 07/934,387

NAME: GRACEY, NANCY J.

REGISTRATION NUMBER: 28,216

REFERENCE/DOCKET NUMBER: 246102002221

TELECOMMUNICATION INFORMATION:

TELEPHONE: 415-813-5600

TELEFAX: 415-494-0792

TELEPAC: 706141

INFORMATION FOR SEQ ID NO: 38:

SEQUENCE CHARACTERISTICS:

LENGTH: 15 base pairs

TYPE: nucleic acid

STRANDEDNESS: single

TOPOLOGY: linear

FEATURE:

NAME/KEY: misc_difference

LOCATION: replace(3..4, "")

OTHER INFORMATION: /note= "This is a formacetal

FEATURE:

NAME/KEY: misc_difference

LOCATION: replace(12..13, "")

OTHER INFORMATION: /note= "This is a formacetal

OTHER INFORMATION: linkage.

US-08-484-192-38

Query Match 100 %; Score 15; DB 1; length 15;

Best Local Similarity 100.0%; Pred. No. 48;

Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

RESULT 14

US-08-484-192-79

Sequence 79, Application US/08484192

; Patent No. 5756291

; GENERAL INFORMATION:

; APPLICANT: GRIFFIN, LINDA C.

; APPLICANT: ALBRECHT, GLENN

; APPLICANT: LATHAM, JOHN

; APPLICANT: LEUNG, LAWRENCE

; APPLICANT: VERMAS, ERIC

; APPLICANT: TOOLE, JOHN J.

; APPLICANT: LATHAM, JOHN

; APPLICANT: LEUNG, LAWRENCE

; APPLICANT: VERMAS, ERIC

; APPLICANT: TOOLE, JOHN J.

TITLE OF INVENTION: APTAMERS SPECIFIC FOR BIOMOLECULES AND

NUMBER OF SEQUENCES: 181

CORRESPONDENCE ADDRESS: MORRISON & FOERSTER

ADRESSE: MORRISON & FOERSTER

STREET: 755 PAGE MILL ROAD

CITY: PALO ALTO

STATE: CALIFORNIA

COUNTRY: USA

ZIP: 94304

COMPUTER READABLE FORM:

COMPUTER: IBM PC compatible

OPERATING SYSTEM: PC-DOS/MS-DOS

SOFTWARE: PatentIn Release #1.0, Version #1.25

CURRENT APPLICATION DATA:

APPLICATION NUMBER: US/08/484,192

FILING DATE: 21-AUG-1992

CLASSIFICATION: 435

PRIORITY APPLICATION DATA:

APPLICATION NUMBER: US 07/934,387

NAME: GRACEY, NANCY J.

REGISTRATION NUMBER: 28,216

REFERENCE/DOCKET NUMBER: 246102002221

TELECOMMUNICATION INFORMATION:

TELEPHONE: 415-813-5600

TELEFAX: 415-494-0792

TELEPAC: 706141

INFORMATION FOR SEQ ID NO: 39:

SEQUENCE CHARACTERISTICS:

LENGTH: 15 base pairs

TYPE: nucleic acid

STRANDEDNESS: single

TOPOLOGY: linear

FEATURE:

NAME/KEY: misc_difference

LOCATION: replace(3..4, "")

OTHER INFORMATION: /note= "This is a formacetal

FEATURE:

NAME/KEY: misc_difference

LOCATION: replace(12..13, "")

OTHER INFORMATION: /note= "This is a formacetal

OTHER INFORMATION: linkage.

US-08-484-192-39

Query Match 100 %; Score 15; DB 1; length 15;

Best Local Similarity 100.0%; Pred. No. 48;

Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

RESULT 15

US-08-484-192-40

Sequence 40, Application US/08484192

; Patent No. 5756291

; GENERAL INFORMATION:

; APPLICANT: GRIFFIN, LINDA C.

; APPLICANT: ALBRECHT, GLENN

; APPLICANT: LATHAM, JOHN

; APPLICANT: LEUNG, LAWRENCE

; APPLICANT: VERMAS, ERIC

; APPLICANT: TOOLE, JOHN J.

; APPLICANT: LATHAM, JOHN

; APPLICANT: LEUNG, LAWRENCE

; APPLICANT: VERMAS, ERIC

; APPLICANT: TOOLE, JOHN J.

TITLE OF INVENTION: APTAMERS SPECIFIC FOR BIOMOLECULES AND

NUMBER OF SEQUENCES: 181

CORRESPONDENCE ADDRESS: MORRISON & FOERSTER

ADRESSE: MORRISON & FOERSTER

